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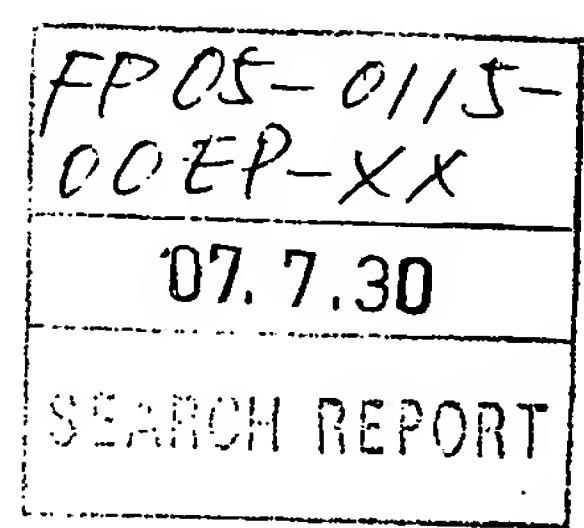
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(54) **MEDICINAL COMPOSITION**

(57) A pharmaceutical composition comprising: an active ingredient consisting of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinoliniccarboxamide, salt thereof, or solvate of the foregoing; and (i) a compound, a 5% (w/w) aqueous solution or suspension of which has a pH of 8 or more, and/or (ii) silicic acid, salt thereof, or solvate of the foregoing is a highly

stable pharmaceutical composition, wherein under humidified and heated storage conditions, the decomposition of said compound is sufficiently reduced, or the gelation on the surface of the pharmaceutical composition is sufficiently inhibited.

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Description

Technical Field

5 [0001] The present invention relates to pharmaceutical compositions.

Background Art

10 [0002] Nitrogen-containing aromatic ring derivatives disclosed in Patent Document 1 have actions in vitro such as 1) the inhibition of infiltrating tube formation by vascular endothelial cells induced by an angiogenic factor mixture solution; 2) the inhibition of tube formation by vascular endothelial cells induced specifically by a single angiogenic factor; 3) the inhibition of angiogenic factor receptor kinase; and 4) the inhibition of cancer cell growth, and hence are extremely useful as angiogenic inhibitors and the like.

15 [0003] [Patent Document 1]: WO 02/32872

Disclosure of the Invention

Problems to be Solved by the Invention

20 [0004] The present inventors have found that while studying on formulating the above nitrogen-containing aromatic ring derivatives, pharmaceutical compositions containing the above nitrogen-containing aromatic ring derivatives as active ingredients are sometimes rendered unstable. Specifically, of the nitrogen-containing aromatic ring derivatives disclosed in Patent Document 1, this holds true for the nitrogen-containing aromatic ring derivatives having a structure wherein the quinoline skeleton is linked to another heterocyclic group through an ether bond. In particular, in pharmaceutical compositions, such nitrogen-containing aromatic ring derivatives are readily decomposed under humidified and heated storage conditions; and moreover, gelation readily occurs on the surface of the pharmaceutical compositions, so that when the pharmaceutical compositions are stored under humidified conditions, delayed dissolution of the active ingredients may occur due to moisture absorption.

25 [0005] Accordingly, an object of the present invention is to provide a stable pharmaceutical composition comprising a nitrogen-containing aromatic ring derivative, wherein under humidified and heated storage conditions, the decomposition of the above derivative is sufficiently reduced, or the gelation on the surface of the pharmaceutical composition is sufficiently inhibited.

Means for Solving the Problems

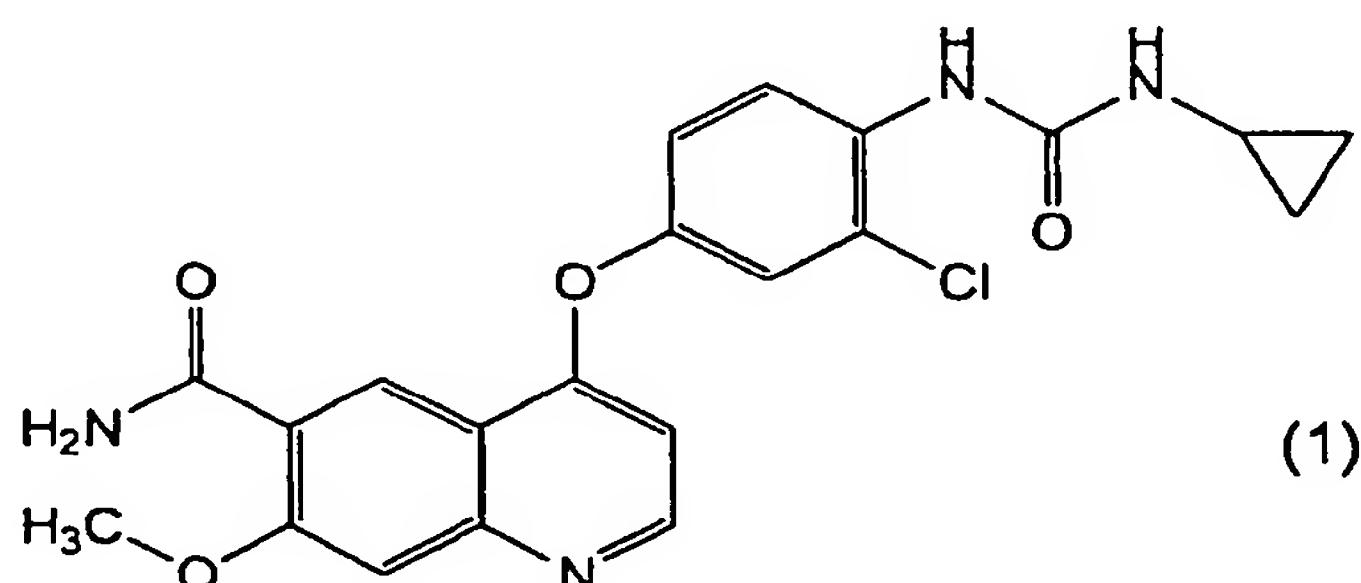
30 [0006] To attain the above object, the present invention provides the pharmaceutical composition described below.

35 [0007] The pharmaceutical composition comprising:

40 an active ingredient consisting of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinoline-carboxamide represented by Formula (1) described below, salt thereof, or solvate of the foregoing; and

- (i) a compound, a 5% (w/w) aqueous solution or suspension of which has a pH of 8 or more; and/or
- (ii) silicic acid, salt thereof, or solvate of the foregoing.

45 [Formula 1]

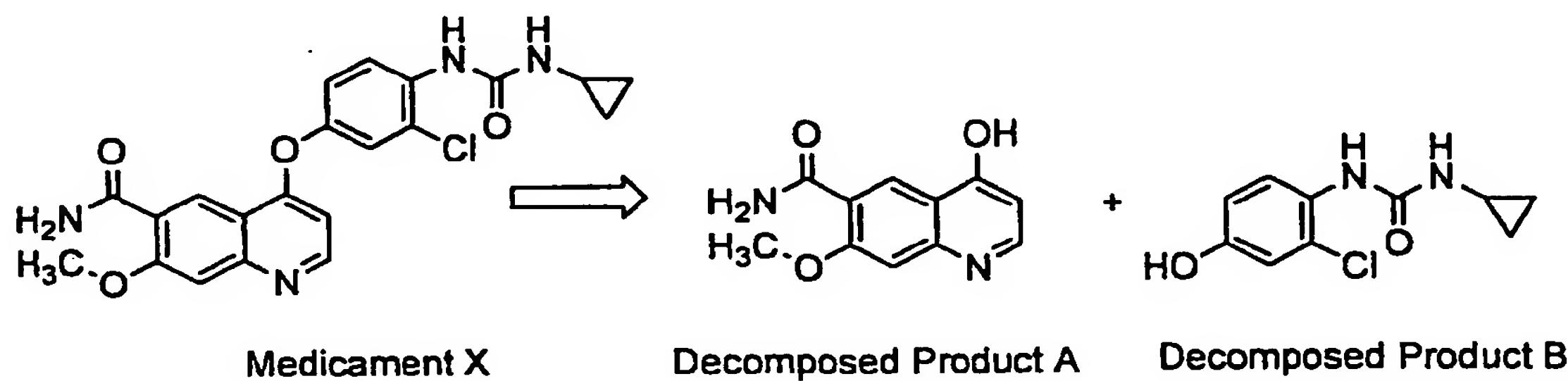


[0008] In such pharmaceutical composition, the decomposition of the compound represented by Formula (1), an active ingredient, under humidified and heated storage conditions is sufficiently reduced. Moreover, the gelation on the surface of the pharmaceutical composition is inhibited, and thereby the problem of delayed dissolution of the active ingredient after the pharmaceutical composition has been kept under humidified storage conditions is solved. Therefore, difficulties during a disintegration test or a dissolution test caused by the surface gelation of the pharmaceutical composition are eliminated, and humidity and the like do not affect the pharmaceutical composition so that the quality of the pharmaceutical composition can be ensured for a long time.

[0009] It is considered that under humidified and heated storage conditions, the suppression of the decomposition is brought about mainly by (i) the compound whose pH of a 5% (w/w) aqueous solution or suspension thereof is 8 or more, whereas the inhibition of the gelation is brought about mainly by (ii) silicic acid, salt thereof, or solvate of the foregoing. Therefore, according to requirements for the pharmaceutical composition, (i) or (ii) can be added alone or in combination thereto.

[0010] Further, it is considered that the decomposition of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide (hereinafter, also referred to as the "medicament X"), salt thereof, or solvate of the foregoing under humidified and heated storage conditions proceed based on the following mechanism (hereinafter, also the decomposed product having a quinoline skeleton is referred to as the "decomposed product A" and the decomposed product having 3-chloro-4-(cyclopropylaminocarbonyl)amino group as the "decomposed product B").

[Formula 2]



[0011] According to these findings, a process for improving stability of the pharmaceutical composition and a process for inhibiting gelation thereof are provided. Specifically, provided are a process for improving stability of the pharmaceutical composition comprising an active ingredient consisting of the compound represented by Formula (1) described above, salt thereof, or solvate of the foregoing by the process of adding the compound whose pH of a 5% (w/w) aqueous solution or suspension thereof is 8 or more, and a process for inhibiting gelation of the pharmaceutical composition comprising an active ingredient consisting of the compound represented by Formula (1) described above, salt thereof, or solvate of the foregoing by the process of adding silicic acid, salt thereof, or solvate of the foregoing.

[0012] In the present invention, (i) the compound whose pH of a 5% (w/w) aqueous solution or suspension thereof is 8 or more is preferably one or more selected from the group consisting of magnesium oxide, calcium oxide, sodium carbonate, disodium hydrogenphosphate, sodium citrate, dipotassium hydrogenphosphate, sodium acetate, sodium hydrogencarbonate, and sodium hydroxide; and (ii) silicic acid, salt thereof, or solvate of the foregoing is preferably one or more selected from the group consisting of light anhydrous silicic acid, silicon dioxide hydrate, calcium silicate, magnesium silicate, magnesium aluminosilicate, magnesium aluminometasilicate, magnesium aluminum silicate, synthetic aluminum silicate, and hydrous silicic dioxide.

Effect of the Invention

[0013] Provided is the highly stable pharmaceutical composition comprising a nitrogen-containing aromatic ring derivative, wherein under humidified and heated storage conditions, the decomposition of the above derivative is sufficiently reduced, or the gelation on the surface of the pharmaceutical composition is sufficiently inhibited.

Brief Description of the Drawings

55 [0014]

Figure 1 illustrates the relationship between pH and the decomposed product A; Figure 2 illustrates the dissolution test results for the Example 2;

Figure 3 illustrates the dissolution test results for the Example 3;
 Figure 4 illustrates the dissolution test results for the Example 4 and for the Comparative Example 2;
 Figure 5 illustrates the dissolution test results for the Examples 5, 6 and 7; and
 Figure 6 is a graph illustrating the amount of the decomposed product A generated when various kinds of stabilizers
 5 were added at various concentrations.

Best Mode for Carrying Out the Invention

[0015] An embodiment of the present invention is explained in detail in the following paragraphs. Herein, the expression "having a diffraction peak at a diffraction angle $(2\theta \pm 0.2^\circ)$ of X° " means to have a diffraction peak at a diffraction angle (2θ) of from $(X - 0.2)^\circ$ to $(X + 0.2)^\circ$. In general, a diffraction angle (2θ) in a powder X-ray diffraction has an error within a range of $\pm 0.2^\circ$, and hence it should be understood that the values of the diffraction angles may include numerals on the order of $\pm 0.2^\circ$. Accordingly, the present invention encompasses not only crystals having completely matching diffraction angles of the peaks in a powder X-ray diffraction, but also crystals having matching diffraction angles of the peaks within the errors of about $\pm 0.2^\circ$.

(Active Ingredient)

[0016] The pharmaceutical composition in accordance with the present invention comprises the compound represented by Formula (1), salt thereof, or solvate of the foregoing as an active ingredient. The active ingredient represented by Formula (1) may be the polymorphic crystals (A') or the polymorphic crystals (B') described below.

[0017] As the polymorphic crystals (A'), the polymorphic crystals (A') of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide having a diffraction peak at a diffraction angle $(2\theta \pm 0.2^\circ)$ of 15.75° in a powder X-ray diffraction can be employed. This polymorphic crystals (A') may also have diffraction peaks at diffraction angles $(2\theta \pm 0.2^\circ)$ of 9.98° and 11.01° in a powder X-ray diffraction.

[0018] In an infrared spectrum (potassium bromide), these polymorphic crystals (A') may preferably have absorbance at $3452.3 \pm 2.5 \text{ cm}^{-1}$ and also at $1712.2 \pm 1.0 \text{ cm}^{-1}$.

[0019] As the polymorphic crystals (B'), the polymorphic crystals (B') of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide having a diffraction peak at a diffraction angle $(2\theta \pm 0.2^\circ)$ of 21.75° in a powder X-ray diffraction can be employed. This polymorphic crystals (B') may also have diffraction peaks at diffraction angles $(2\theta \pm 0.2^\circ)$ of 12.43° and 16.56° in a powder X-ray diffraction.

[0020] In an infrared spectrum (potassium bromide), these polymorphic crystals (B'), active ingredients, may preferably have absorbance at $1557.6 \pm 1.0 \text{ cm}^{-1}$ and also at $1464.4 \pm 1.0 \text{ cm}^{-1}$.

[0021] It is particularly preferred that the active ingredient represented by Formula (1) is salts, solvates, or crystals of these described below.

[0022] In particular, a suitable active ingredient is crystals of hydrochloride, hydrobromide, p-toluenesulfonate, sulfate, methanesulfonate, or ethanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide or of solvate thereof.

[0023] Specifically, crystals of methanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide or solvate thereof; crystals of ethanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide or solvate thereof; crystals of methanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide; crystals of hydrate of methanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide; crystals of dimethyl sulfoxide solvate of methanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide; crystals of acetic acid solvate of methanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide; crystals of ethanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide; and crystals of dimethyl sulfoxide solvate of ethanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide are suitable.

[0024] The crystals of methanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide are preferably the crystals (A), the crystals (B), or the crystals (C) described below.

[0025] Specifically, the crystals (A) having diffraction peaks at diffraction angles $(2\theta \pm 0.2^\circ)$ of 9.65° and 18.37° in a powder X-ray diffraction; the crystals (B) having diffraction peaks at diffraction angles $(2\theta \pm 0.2^\circ)$ of 5.72° and 13.84° in a powder X-ray diffraction; and the crystals (C) having diffraction peaks at diffraction angles $(2\theta \pm 0.2^\circ)$ of 14.20° and 17.59° in a powder X-ray diffraction are preferred.

[0026] Further, the crystals of hydrate of methanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide are preferably the crystals (F) having diffraction peaks at diffraction angles $(2\theta \pm 0.2^\circ)$ of 8.02° and 18.14° in a powder X-ray diffraction; and the crystals of acetic acid solvate of methanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide are preferably the crys-

tals (I) having diffraction peaks at diffraction angles ($2\theta \pm 0.2^\circ$) of 9.36° and 12.40° in a powder X-ray diffraction.

[0027] Moreover, the crystals of ethanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide are preferably the crystals (α) having diffraction peaks at diffraction angles ($2\theta \pm 0.2^\circ$) of 15.70° and 17.18° in a powder X-ray diffraction; and the crystals of ethanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide are preferably the crystals (β) having diffraction peaks at diffraction angles ($2\theta \pm 0.2^\circ$) of 6.48° and 9.58° in a powder X-ray diffraction.

(Process for Preparing the Active Ingredient)

[0028] As for the process for preparing the compound represented by Formula (1), the description in WO 02/32872 can be used as a reference. The processes for preparing the polymorphic crystals (A') and the polymorphic crystals (B') are described in the following paragraphs.

[0029] The polymorphic crystals (A') can be obtained by a preparation process described below: 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide is dissolved in an organic solvent, a good solvent (for example, dimethyl sulfoxide, dimethylimidazolidine, 1-methyl-2-pyrrolidinone, N,N-dimethylformamide, N,N-dimethylacetamide, acetic acid, sulfolane, etc.), and then thereto a poor solvent (for example, water, acetone, acetonitrile, ethyl acetate, isopropyl acetate, methanol, ethanol, n-propanol, isopropanol, or a mixture thereof, etc.) is admixed rapidly (for example, within 10 min.).

[0030] The polymorphic crystals (A') can be obtained by another preparation process described below: 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide is dissolved while stirring in an organic solvent, a good solvent (for example, dimethyl sulfoxide, dimethylimidazolidine, 1-methyl-2-pyrrolidinone, N,N-dimethylformamide, N,N-dimethylacetamide, acetic acid, sulfolane, etc.), and then thereto a poor solvent (for example, water, acetone, acetonitrile, ethyl acetate, isopropyl acetate, methanol, ethanol, n-propanol, isopropanol, or a mixture thereof, etc.) is admixed so that the resultant crystals precipitate when the stirring is stopped.

[0031] The polymorphic crystals (A') can be obtained by still another preparation process described below: 7-methoxy-4-chloroquinoline-6-carboxamide and 1-(2-chloro-4-hydroxyphenyl)-3-cyclopropylurea are reacted in the presence of a base (for example, potassium t-butoxide, cesium carbonate, potassium carbonate, etc.) in an organic solvent that works a good solvent for 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide (for example, dimethyl sulfoxide (DMSO), dimethylimidazolidine, 1-methyl-2-pyrrolidinone, N,N-dimethylformamide, N,N-dimethylacetamide, sulfolane, etc.), and then thereto a poor solvent is admixed rapidly (for example, within 10 min.).

[0032] More specifically, for example, to a mixture of 1-(2-chloro-4-hydroxyphenyl)-3-cyclopropylurea, 7-methoxy-4-chloroquinoline-6-carboxamide (one equivalent or more based on 1-(2-chloro-4-hydroxyphenyl)-3-cyclopropylurea), and potassium t-butoxide (one equivalent or more based on 1-(2-chloro-4-hydroxyphenyl)-3-cyclopropylurea), DMSO in a volume 5 to 10 times based on 1-(2-chloro-4-hydroxyphenyl)-3-cyclopropylurea is added at room temperature, and then the mixture is heated to from 55 to 75°C while stirring for 20 hr. or longer to allow the reaction to proceed. To this reaction mixture, while heating at from 60 to 65°C and stirring, a poor solvent (20 to 50% acetone in water or 20 to 50% 2-propanol in water) in a volume 15 times based on 1-(2-chloro-4-hydroxyphenyl)-3-cyclopropylurea is introduced within 8 min. so that crystals can appear. Further, it is preferred to add the seed crystals when the poor solvent is introduced to allow the crystals to appear. The polymorphic crystals (A') can be obtained by stirring the resulting reaction mixture, in which the crystals appeared, at the temperature ranging from room temperature to 40°C generated by heating for 3 hr. or longer to collect the crystals by filtration.

[0033] The polymorphic crystals (B') can be obtained by a preparation process described below: 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide is dissolved in an organic solvent, a good solvent (for example, DMSO, dimethylimidazolidine, 1-methyl-2-pyrrolidinone, N,N-dimethylformamide, N,N-dimethylacetamide, acetic acid, sulfolane, etc.), and then thereto a poor solvent (for example, water, acetone, acetonitrile, ethyl acetate, isopropyl acetate, methanol, ethanol, n-propanol, isopropanol, or a mixture thereof, etc.) is admixed slowly (for example, for 1 hr. or longer). When the poor solvent is admixed slowly, crystals appeared, but when stirring is stopped, the resultant crystals spreads all over the solvent.

[0034] More specifically, for example, to 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide, a good solvent (DMSO or 1-methyl-2-pyrrolidinone) in a volume from 4 to 5 times based on 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide is added, and then the mixture is heated to 80°C or higher while stirring to dissolve the solute. To this mixture, while heating at from 65 to 85°C and stirring, a poor solvent (isopropyl acetate, ethyl acetate, methanol, or isopropanol) in a volume from 10 to 20 times based on 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide is introduced over 30 min. or longer so that crystals can appear. Further, it is preferred to add the seed crystals when the poor solvent is introduced to allow the crystals to appear. The polymorphic crystals (B') can be obtained by stirring the resulting reaction mixture, in which the crystals appeared, while heating at 70°C or higher for 30 min. or longer, and further by stirring at room temperature to collect the crystals by filtration.

[0035] The polymorphic crystals (B') can be obtained by another preparation process described below: 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide is dissolved while stirring in an organic solvent, a good solvent (for example, DMSO, dimethylimidazolidine, 1-methyl-2-pyrrolidinone, N,N-dimethylformamide, N,N-dimethylacetamide, acetic acid, sulfolane, etc.), and then a poor solvent (for example, water, acetone, acetonitrile, ethyl acetate, isopropyl acetate, methanol, ethanol, n-propanol, isopropanol, or a mixture thereof, etc.) is admixed so that when stirring is stopped, the resultant crystals spreads all over the solvent.

[0036] The polymorphic crystals (B') can be obtained by still another preparation process described below: 7-methoxy-4-chloroquinoline-6-carboxamide and 1-(2-chloro-4-hydroxyphenyl)-3-cyclopropylurea are reacted in the presence of a base (for example, potassium t-butoxide, cesium carbonate, potassium carbonate, etc.) in an organic solvent that works a good solvent for 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide (for example, DMSO, dimethylimidazolidinone, 1-methyl-2-pyrrolidinone, N,N-dimethylformamide, N,N-dimethylacetamide, sulfolane, etc.), and then thereto a poor solvent (for example, water, acetone, acetonitrile, ethyl acetate, isopropyl acetate, methanol, ethanol, n-propanol, isopropanol, or a mixture thereof, etc.) is admixed slowly (for example, for 30 min. or longer).

[0037] More specifically, for example, to a mixture of 1-(2-chloro-4-hydroxyphenyl)-3-cyclopropylurea, 7-methoxy-4-chloroquinoline-6-carboxamide (one equivalent or more based on 1-(2-chloro-4-hydroxyphenyl)-3-cyclopropylurea), and potassium t-butoxide (one equivalent or more based on 1-(2-chloro-4-hydroxyphenyl)-3-cyclopropylurea), DMSO in a volume from 5 to 10 times based on 1-(2-chloro-4-hydroxyphenyl)-3-cyclopropylurea is added at room temperature, and then the mixture is heated to from 55 to 75°C while stirring for 20 hr. or longer to allow the reaction to proceed. To this reaction mixture, while heating at from 60 to 65°C and stirring, a poor solvent (33% acetone in water) in a volume 15 times based on 1-(2-chloro-4-hydroxyphenyl)-3-cyclopropylurea is introduced over 2 hr. or longer so that crystals can appear. The polymorphic crystals (B') can be obtained by stirring the resulting reaction mixture, in which the crystals appeared, while heating at 40°C for 3 hr. or longer to collect the crystals by filtration.

[0038] The polymorphic crystals (B') can be obtained by still another preparation process described below: the polymorphic crystals (A') of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide having a diffraction peak at a diffraction angle ($2\theta \pm 0.2^\circ$) of 15.75° in a powder X-ray diffraction is suspended and heated in a mixed solution of an organic solvent which is a good solvent for the above polymorphic crystals and a poor solvent for the above polymorphic crystals. It is preferred that the polymorphic crystals (A') used for this purpose also has diffraction peaks at diffraction angles ($2\theta \pm 0.2^\circ$) of 9.98° and 11.01° .

[0039] The polymorphic crystals (B') can be obtained by still another preparation process described below: the polymorphic crystals (A') of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide having absorbance at a wavenumber of $3452.3 \pm 2.5 \text{ cm}^{-1}$ in an infrared absorption spectrum (potassium bromide) is suspended and heated in a mixed solution of a good solvent for the above polymorphic crystals and a poor solvent for the above polymorphic crystals. It is preferred that the polymorphic crystals (A') used for this purpose has absorbance at a wavenumber of $3452.3 \pm 2.5 \text{ cm}^{-1}$ (and also at $1712.2 \pm 1.0 \text{ cm}^{-1}$) in an infrared absorption spectrum (potassium bromide).

[0040] The crystals of hydrochloride or hydrobromide of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide; the crystals of p-toluenesulfonate or sulfate of the same; the crystals (A) of the same; the crystals (B) of the same; the crystals (C) of the same; the crystals of dimethyl sulfoxide solvate of methanesulfonate of the same; the crystals (F) of the same; the crystals (I) of the same; the crystals (α) of the same; the crystals (β) of the same; and the crystals of dimethyl sulfoxide solvate of ethanesulfonate of the same can be obtained by preparation processes described in the following paragraphs.

[0041] The crystals of hydrochloride or hydrobromide can be obtained by mixing the carboxamide and a solvent to dissolve the carboxamide therein, to which hydrochloric acid or hydrobromic acid is then added. More specifically, for example, the carboxamide and the solvent are mixed to dissolve the carboxamide therein by heating, and then thereto hydrochloric acid or hydrobromic acid is added followed by gradually cooling the resulting mixture to room temperature so that the crystals of hydrochloride or hydrobromide can be prepared. As a solvent, an alcohol such as methanol, ethanol, 1-propanol, and 2-propanol can be used, and ethanol is preferred. Further, water may be optionally added to the alcohol. There is no particular restriction on the amount of the solvent, but the solvent preferably from 10 to 30 times, more preferably 20 times, as much as the solute is used. As for the amount of hydrochloric acid or hydrobromic acid, from 1.0 to 1.5 equivalents, preferably 1.1 equivalents, based on the solute can be used. There is no particular restriction on the heating temperature, but the heating temperature is preferably from 60°C to the reflux temperature, and more preferably the reflux temperature. For cooling, it may take from 10 min. to 24 hr. to cool down gradually from the heating temperature to room temperature.

[0042] The crystals of a p-toluenesulfonate or sulfate can be obtained by mixing the carboxamide, a solvent, and p-toluenesulfonic acid or sulfuric acid to dissolve the carboxamide therein. More specifically, for example, the carboxamide, the solvent, and p-toluenesulfonic acid or sulfuric acid are mixed to dissolve the carboxamide therein by heating followed by gradually cooling the resulting mixture to room temperature so that the crystals of p-toluenesulfonate or sulfate can

be prepared. As a solvent, for example, dimethyl sulfoxide, N,N-dimethylformamide, N,N-dimethylacetamide, etc., can be used, and dimethyl sulfoxide is preferred. There is no particular restriction on the amount of the solvent, but the solvent preferably from 10 to 30 times, more preferably 20 times, as much as the solute is used. As for the amount of p-toluenesulfonic acid or sulfuric acid, from 1.0 to 1.5 equivalents, preferably 1.2 equivalents, based on the solute can be used. There is no particular restriction on the heating temperature, but the heating temperature is preferably from 60°C to the reflux temperature, more preferably from 70 to 100°C, and still more preferably 80°C. For cooling, it may take from 10 min. to 24 hr. to cool down gradually from the heating temperature to room temperature.

[0043] The crystals (A) can be obtained by a preparation process comprising mixing 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide, a solvent, and methanesulfonic acid to dissolve. More specifically, for example, the carboxamide, the solvent, and methanesulfonic acid are mixed to dissolve the carboxamide therein by heating followed by gradually cooling the resulting mixture to room temperature so that the crystals (A) of methanesulfonate can be prepared. As a solvent, for example, methanol, ethanol, 2-propanol, etc., can be used, and methanol is preferred. There is no particular restriction on the amount of the solvent, but the solvent preferably from 10 to 30 times, more preferably 20 times, as much as the solute is used. As for the amount of methanesulfonic acid, from 1.0 to 1.5 equivalents, preferably 1.2 equivalents, based on the solute can be used. There is no particular restriction on the heating temperature, but the heating temperature is preferably from 60°C to the reflux temperature, and more preferably from 70 to 80°C. For cooling, it may take from 1 to 24 hr., preferably from 3 to 12 hr., to cool down gradually from the heating temperature to room temperature.

[0044] The crystals (A) can be also obtained by a preparation process comprising mixing 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide, acetic acid, and methanesulfonic acid to dissolve. More specifically, for example, the carboxamide, acetic acid, and methanesulfonic acid are mixed to dissolve the carboxamide therein by heating, and then thereto a poor solvent is added followed by gradually cooling the resulting mixture to room temperature so that the crystals (A) of methanesulfonate can be prepared. Further, it is preferred to add the seed crystals (A) of methanesulfonate with the poor solvent. There is no particular restriction on the amount of acetic acid, but acetic acid preferably from 5 to 20 times, more preferably 10 times, as much as the solute is used. As for the amount of methanesulfonic acid, from 1.0 to 2.5 equivalents, preferably from 1.4 to 2.2 equivalents, based on the solute can be used. As a poor solvent, for example, methanol, ethanol, etc., can be used, and ethanol is preferred. There is no particular restriction on the amount of the poor solvent, but the poor solvent preferably from 10 to 30 times, more preferably 20 times, as much as the solute is used. Further, the poor solvent can be added at once, or divided into from 2 to 4 parts, preferably 2 parts, for addition. In such case, the amount ratio of the first addition and the second addition is from 1:1 to 3:1, and preferably 3:2. There is no particular restriction on the heating temperature, but the heating temperature is preferably from 50°C to the reflux temperature, and more preferably 50°C. For cooling, it may take from 10 min. to 6 hr., preferably from 1 to 2 hr., to cool down gradually from the heating temperature to room temperature.

[0045] The crystals (B) can be obtained by a preparation process comprising drying (for example, by drying under aeration) the crystals (I) of acetic acid solvate of methanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide to remove acetic acid.

[0046] The crystals (C) can be obtained by a preparation process comprising heating crystals of dimethyl sulfoxide solvate of methanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide (and preferably cooling down gradually to room temperature). This preparation process can be conducted either in the presence of or in the absence of a solvent. When a solvent is used, as the solvent, for example, ethyl acetate, isopropyl acetate, n-butyl acetate, etc., can be used, and n-butyl acetate is preferred. There is no particular restriction on the heating temperature, but the heating temperature is preferably from 70°C to the reflux temperature, and more preferably the reflux temperature.

[0047] The crystals (C) can be also obtained by a preparation process comprising mixing the crystals (I) of acetic acid solvate of methanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide and a solvent. In this preparation process, as a solvent, for example, an alcohol such as methanol, ethanol, and 2-propanol can be used, and ethanol is preferred. There is no particular restriction on the stirring temperature, but the stirring temperature is preferably from 20 to 60°C, and more preferably 40°C.

[0048] The crystals (C) can be still also obtained by a preparation process comprising mixing 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide, acetic acid, and methanesulfonic acid to dissolve. More specifically, for example, the carboxamide, acetic acid, and methanesulfonic acid are mixed to dissolve the carboxamide therein by heating, and then thereto 2-propanol as a poor solvent is added followed by gradually cooling the resulting mixture to about 15°C so that the crystals (C) of methanesulfonate can be prepared. Further, it is preferred to add the seed crystals (C) of methanesulfonate with the poor solvent and to add isopropyl acetate to accelerate the appearance of the crystals. There is no particular restriction on the amount of acetic acid, but acetic acid preferably from 5 to 10 times, more preferably from 7 to 8 times, as much as the solute is used. As for the amount of methanesulfonic acid, from 1.0 to 1.5 equivalents, preferably 1.2 equivalents, based on the solute can be used. There is no particular restriction on the amount of the poor solvent, but the poor solvent preferably from 2 to 10 times, more preferably from

4 to 5 times, as much as the solute is used. When isopropyl acetate is added, there is no particular restriction on the amount of isopropyl acetate, but isopropyl acetate preferably from 2 to 10 times, more preferably 5 times, as much as the solute is used. There is no particular restriction on the heating temperature, but the heating temperature is preferably 40°C. For cooling, it may take from 10 min. to 6 hr., preferably from 1 to 2 hr., to cool down gradually from the heating temperature to about 15°C.

[0049] In another preparation process wherein 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide, acetic acid, and methanesulfonic acid are mixed to dissolve, the carboxamide, acetic acid, and methanesulfonic acid are mixed to dissolve the carboxamide therein at room temperature (or about 30°C), and then thereto 2-propanol as a poor solvent is added followed by gradually cooling the resulting mixture to about 15°C. Resultant crystals are collected by filtration, and then the above crystals and a solvent are mixed and stirred so that the crystals (C) of methanesulfonate can be prepared. Further, it is preferred to add the seed crystals (C) of methanesulfonate with the poor solvent. There is no particular restriction on the amount of acetic acid, but acetic acid preferably from 5 to 20 times, more preferably 10 times, as much as the solute is used. As for the amount of methanesulfonic acid, from 1.0 to 2.5 equivalents, preferably from 1.8 to 2.2 equivalents, based on the solute can be used. There is no particular restriction on the amount of the poor solvent, but the poor solvent preferably from 10 to 30 times, more preferably 20 times, as much as the solute is used. For cooling, it may take from 10 min. to 4 hr., preferably from 30 min. to 2 hr., to cool down gradually from room temperature (or about 30°C) to about 15°C. As a solvent to be mixed with the collected crystals, for example, an alcohol such as methanol, ethanol, and 2-propanol can be used, and ethanol is preferred.

[0050] In further still another process for preparing the crystals (C), the crystals (B) of methanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide is moisturized.

[0051] Crystals of dimethyl sulfoxide solvate of methanesulfonate can be obtained by mixing the carboxamide, dimethyl sulfoxide, and methanesulfonic acid to dissolve the carboxamide therein by heating, and then by adding thereto a poor solvent followed by cooling the resulting mixture to about 15°C. Further, it is preferred to add the seed crystals (A) of methanesulfonate with the poor solvent. There is no particular restriction on the amount of dimethyl sulfoxide, but dimethyl sulfoxide preferably from 5 to 20 times, more preferably from 8 to 10 times, as much as the solute is used. As for the amount of methanesulfonic acid, from 1.0 to 4.0 equivalents, preferably from 1.2 to 3.5 equivalents, based on the solute can be used. As a poor solvent, for example, ethyl acetate, isopropyl acetate, 1-propanol, 2-propanol, etc., can be used, and ethyl acetate and 2-propanol are preferred. There is no particular restriction on the amount of the poor solvent, but the poor solvent preferably from 10 to 30 times, more preferably 20 times, as much as the solute is used. Further, the poor solvent can be added at once, or divided into from 2 to 4 parts, preferably 2 parts, for addition. In such case, the amount ratio of the first addition and the second addition is from 1:1 to 1:5, and preferably 1:4. There is no particular restriction on the heating temperature, but the heating temperature is preferably from 50 to 100°C, and more preferably 60 to 80°C. For cooling, it may take from 10 min. to 6 hr., preferably from 1 to 2 hr., to cool down from the heating temperature to about 15°C.

[0052] The crystals (F) can be obtained by a preparation process comprising mixing 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide, acetic acid, and methanesulfonic acid to dissolve. More specifically, for example, the carboxamide, acetic acid, and methanesulfonic acid are mixed to dissolve the carboxamide therein by heating, and then thereto a poor solvent is added followed by gradually cooling the resulting mixture to room temperature so that the crystals (F) of hydrate of methanesulfonate can be prepared. Further, it is preferred to add the seed crystals (A) of methanesulfonate with the poor solvent. There is no particular restriction on the amount of acetic acid, but acetic acid preferably from 5 to 20 times, more preferably 10 times, as much as the solute is used. As for the amount of methanesulfonic acid, from 1.0 to 2.0 equivalents, preferably from 1.3 to 1.6 equivalents, based on the solute can be used. As a poor solvent, for example, ethyl acetate and isopropyl acetate can be used, and ethyl acetate is preferred. There is no particular restriction on the amount of the poor solvent, but the poor solvent preferably from 10 to 30 times, more preferably 20 times, as much as the solute is used. Further, the poor solvent can be added at once, or divided into from 2 to 4 parts, preferably 2 parts, for addition. In such case, the amount ratio of the first addition and the second addition is from 1:1 to 1:5, and preferably 1:3. There is no particular restriction on the heating temperature, but the heating temperature is preferably from 40 to 60°C, and more preferably 50°C. For cooling, it may take from 10 min. to 6 hr., preferably from 2 to 4 hr., to cool down gradually from the heating temperature to room temperature.

[0053] The crystals (I) can be obtained by a preparation process comprising mixing 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide, acetic acid, and methanesulfonic acid to dissolve. More specifically, for example, the carboxamide, acetic acid, and methanesulfonic acid are mixed to dissolve the carboxamide therein by heating, and then thereto a poor solvent is added followed by gradually cooling the resulting mixture to room temperature so that the crystals (I) of an acetic acid solvate of methanesulfonate can be prepared. Further, it is preferred to add the seed crystals (C) of methanesulfonate with the poor solvent and to add isopropyl acetate to accelerate the appearance of the crystals. There is no particular restriction on the amount of acetic acid, but acetic acid preferably from 5 to 10 times, more preferably from 7 to 8 times, as much as the solute is used. As for the amount of methanesulfonic acid, from 1.0 to 1.5 equivalents, preferably 1.2 equivalents, based on the solute can be used. As a poor solvent, for

example, 1-propanol 1-butanol, tert-butanol, etc., can be used, and 1-propanol is preferred. There is no particular restriction on the amount of the poor solvent, but the poor solvent preferably from 5 to 20 times, more preferably from 8 to 10 times, as much as the solute is used. Further, the poor solvent can be added at once, or divided into from 2 to 4 parts, preferably 2 parts, for addition. In such case, the amount ratio of the first addition and the second addition is from 1:1 to 1:5, and preferably 1:3.5. When isopropyl acetate is added, there is no particular restriction on the amount of isopropyl acetate, but isopropyl acetate preferably from 2 to 10 times, more preferably 5 times, as much as the solute is used. There is no particular restriction on the heating temperature, but the heating temperature is preferably 40°C. For cooling, it may take from 10 min. to 6 hr., preferably from 1 to 2 hr., to cool down gradually from the heating temperature to room temperature.

[0054] The crystals (α) can be obtained by a preparation process comprising mixing 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide, a solvent, and ethanesulfonic acid to dissolve. More specifically, for example, the carboxamide, the solvent, and ethanesulfonic acid are mixed to dissolve the carboxamide therein by heating, and then thereto a poor solvent is added followed by gradually cooling the resulting solution to room temperature so that the crystals (α) of ethanesulfonate can be prepared. As a solvent, for example, dimethyl sulfoxide, etc., can be used. There is no particular restriction on the amount of the solvent, but the solvent preferably from 5 to 20 times, more preferably 10 times, as much as the solute is used. As for the amount of ethanesulfonic acid, from 1.0 to 1.5 equivalents, preferably 1.2 equivalents, based on the solute can be used. As a poor solvent, for example, ethyl acetate, etc., can be used. There is no particular restriction on the amount of the poor solvent, but the poor solvent preferably from 5 to 20 times, more preferably 10 times, as much as the solute is used. There is no particular restriction on the heating temperature, but the heating temperature is preferably from 50 to 70°C, and more preferably 60°C. For cooling, it may take from 10 min. to 24 hr., preferably from 1 to 2 hr., to cool down gradually from the heating temperature to room temperature.

[0055] The crystals (β) can be obtained by a preparation process comprising mixing the crystals (α) of ethanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide and a solvent. As a solvent, for example, methanol, ethanol, 2-propanol, etc., can be used, and ethanol is preferred. There is no particular restriction on the amount of the solvent, but the solvent preferably from 5 to 20 times, more preferably 10 times, as much as the solute is used. There is no particular restriction on the amount of water, but water preferably from 1/10 to 1/2 times, more preferably 1/6 times, as much as ethanol is used.

[0056] The crystals (β) can be also obtained by a preparation process comprising mixing 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide, acetic acid, and ethanesulfonic acid to dissolve. More specifically, for example, the carboxamide, acetic acid, and ethanesulfonic acid are mixed to dissolve the carboxamide therein by heating, and then thereto a poor solvent and water are added followed by cooling the resulting mixture to 0°C so that the crystals (β) of a hydrate of ethanesulfonate can be prepared. Further, it is preferred to add the seed crystals (β) of ethanesulfonate with the poor solvent. There is no particular restriction on the amount of acetic acid, but acetic acid preferably from 2.5 to 10 times, more preferably 5 times, as much as the solute is used. As for the amount of ethanesulfonic acid, from 1.0 to 1.5 equivalents, preferably 1.2 equivalents, based on the solute can be used. As a poor solvent, for example, ethanol, 2-propanol, etc., can be used, and 2-propanol is preferred. There is no particular restriction on the amount of the poor solvent, but the poor solvent preferably from 10 to 40 times, more preferably 30 times, as much as the solute is used. Further, the poor solvent can be added at once, or divided into from 2 to 4 parts, preferably 2 parts, for addition. In such case, the amount ratio of the first addition and the second addition is from 1:1 to 1:5, and preferably from 1:1.5 to 1:2. There is no particular restriction on the amount of water, but water preferably from 1/10 to 1/30 times, more preferably 1/20 times, as much as the poor solvent is used. There is no particular restriction on the heating temperature, but the heating temperature is preferably from 50 to 70°C, and more preferably 60°C. For cooling, it may take from 10 min. to 6 hr., preferably from 2 to 4 hr., to cool down from the heating temperature to 0°C.

[0057] Crystals of dimethyl sulfoxide solvate of ethanesulfonate can be obtained by mixing the carboxamide, dimethyl sulfoxide, and ethanesulfonic acid to dissolve the carboxamide therein by heating, and then by adding a poor solvent thereto followed by cooling the resulting solution to 0°C. Further, it is preferred to add the seed crystals (β) of ethanesulfonate with the poor solvent. There is no particular restriction on the amount of dimethyl sulfoxide, but dimethyl sulfoxide preferably from 5 to 20 times, more preferably 10 times, as much as the solute is used. As for the amount of ethanesulfonic acid, from 1.0 to 1.5 equivalents, preferably 1.2 equivalents, based on the solute can be used. As a poor solvent, for example, ethyl acetate, etc., can be used. There is no particular restriction on the amount of the poor solvent, but the poor solvent preferably from 5 to 20 times, more preferably 10 times, as much as the solute is used. Further, the poor solvent can be added at once, or divided into from 2 to 4 parts, preferably 2 parts, for addition. In such case, the amount ratio of the first addition and the second addition is from 1:1 to 3:1, and preferably 3:2. There is no particular restriction on the heating temperature, but the heating temperature is preferably from 50 to 70°C, and more preferably 60°C. For cooling, it may take from 10 min. to 6 hr., preferably from 1 to 2 hr., to cool down from the heating temperature to 0°C.

(Pharmaceutical Composition)

[0058] The pharmaceutical composition in accordance with the present invention comprises: in addition to the active ingredient consisting of the compound represented by Formula (1), salt thereof, or solvate of the foregoing, as described above;

5 (i) a compound, a 5% (w/w) aqueous solution or suspension of which has a pH of 8 or more; and/or
 (ii) silicic acid, salt thereof, or solvate of the foregoing.

10 [0059] Further, the compound whose pH of a 5% (w/w) aqueous solution or suspension thereof is 8 or more contributes to the suppression of the decomposition of the active ingredient under humidified and heated storage conditions, and hence hereinafter is referred to as the "stabilizer." Moreover, silicic acid, salt thereof, or solvate of the foregoing contributes to the inhibition of the gelation of the pharmaceutical composition, and hence hereinafter is referred to as the "gelation inhibitor."

15 [0060] As the stabilizer, magnesium oxide, calcium oxide, sodium carbonate, disodium hydrogenphosphate, sodium citrate, dipotassium hydrogenphosphate, sodium acetate, sodium hydrogencarbonate, and sodium hydroxide are preferred. Of these, magnesium oxide and calcium oxide are particularly preferred in view of an increase in weight and coloration. The amount of the stabilizer to add to the pharmaceutical composition is preferably from 0.5 to 15, more preferably from 1 to 10, and most preferably from 1 to 5 mass parts based on 100 mass parts of the pharmaceutical composition.

20 [0061] As the gelation inhibitor, light anhydrous silicic acid, silicon dioxide hydrate, calcium silicate, magnesium silicate, magnesium aluminosilicate, magnesium aluminometasilicate, magnesium aluminum silicate, synthetic aluminum silicate, and hydrous silicic dioxide are preferred. Of these, light anhydrous silicic acid, silicon dioxide hydrate, and calcium silicate are more preferred, and light anhydrous silicic acid and silicon dioxide hydrate are most preferred. The amount of the gelation inhibitor to add to the pharmaceutical composition is preferably from 4 to 20, and more preferably from 8 to 20 mass parts based on 100 mass parts of the pharmaceutical composition.

25 [0062] In the pharmaceutical composition in accordance with the present invention, in addition to the active ingredient consisting of the compound represented by Formula (1), salt thereof, or solvate of the foregoing, the stabilizer, and the gelation inhibitor, additives such as a diluent, a binder, a lubricant, a disintegrant, a coloring agent, a flavoring agent, an emulsifier, a surfactant, a solubilizing agent, a suspending agent, an isotonizing agent, a buffer, a preservative, an antioxidant, a stabilizing agent, and an absorption promoter can be added thereto.

30 [0063] Examples of diluents include lactose, sucrose, glucose, cornstarch, mannitol, sorbitol, starch, alpha starch, dextrin, crystalline cellulose, light anhydrous silicic acid, aluminum silicate, calcium silicate, magnesium aluminometasilicate, calcium hydrogenphosphate, etc.

35 [0064] Examples of binders include polyvinyl alcohol, methylcellulose, ethylcellulose, gum Arabic, tragacanth, gelatin, shellac, hydroxypropylmethylcellulose, hydroxypropylcellulose, sodium carboxymethylcellulose, polyvinylpyrrolidone, Macrogol, etc.

[0065] Examples of lubricants include magnesium stearate, calcium stearate, sodium stearyl fumarate, talc, polyethylene glycol, colloidal silica, etc.

40 [0066] Examples of disintegrants include crystalline cellulose, agar, gelatin, calcium carbonate, sodium hydrogencarbonate, calcium citrate, dextrin, pectin, low-substituted hydroxypropylcellulose, carboxymethylcellulose, carboxymethylcellulose calcium, croscarmellose sodium, carboxymethyl starch, carboxymethyl starch sodium, carmellose, carmellose sodium, crospovidone, low-substituted carboxymethyl starch sodium, partially alpha starch, etc. The amount of the disintegrant to add to the pharmaceutical composition is preferably from 0.1 to 30, and more preferably from 1 to 20 mass parts based on 100 mass parts of the pharmaceutical composition.

45 [0067] As the disintegrant, low-substituted hydroxypropylcellulose, carboxymethyl starch sodium, carmellose sodium, carmellose calcium, croscarmellose sodium, crospovidone, and partially alpha starch are preferred. Low-substituted hydroxypropylcellulose, carmellose calcium, croscarmellose sodium, crospovidone, and partially alpha starch are more preferred. Croscarmellose sodium is most preferred.

50 [0068] Examples of coloring agents include iron sesquioxide, yellow iron sesquioxide, carmine, caramel, beta-carotene, titanium oxide, talc, riboflavin sodium phosphate, yellow aluminum lake, etc, which have been approved as additives for medicaments.

[0069] Flavoring agents include cocoa powder, menthol, aromatic powder, mentha oil, borneol, powdered cinnamon bark, etc.

55 [0070] Examples of emulsifiers or surfactants include stearyl triethanolamine, sodium lauryl sulfate, lauryl aminopropionic acid, lecithin, glycerin monostearate, sucrose fatty acid ester, glycerin fatty acid ester, etc.

[0071] Examples of solubilizers include polyethylene glycol, propylene glycol, benzyl benzoate, ethanol, cholesterol, triethanolamine, sodium carbonate, sodium citrate, Polysorbate 80, nicotinamide, etc.

[0072] Examples of suspending agents include, in addition to the above surfactants, hydrophilic polymers such as polyvinyl alcohol, polyvinylpyrrolidone, methylcellulose, hydroxymethylcellulose, hydroxyethylcellulose, and hydroxypropylcellulose.

[0073] Examples of isotonizing agent include glucose, sodium chloride, mannitol, sorbitol, etc.

[0074] Examples of buffers include buffer solutions of phosphate, acetate, carbonate, citrate, etc.

[0075] Examples of preservatives include methylparaben, propylparaben, chlorobutanol, benzyl alcohol, phenethyl alcohol, dehydroacetic acid, sorbic acid, etc.

[0076] Examples of antioxidants include sulfite, ascorbic acid, alpha-tocopherol, etc.

[0077] Further, the pharmaceutical composition can be formulated into oral preparations such as tablets, powders, granules, capsules, syrups, troches, and inhalants; external preparations such as suppositories, ointments, ophthalmic ointments, tapes, eye drops, nasal drops, ear drops, cataplasms, and lotions; or injections. Oral preparations are formulated by combining the above additives as desired. Moreover, optionally surface of these oral preparations may be coated.

[0078] External preparations are formulated by combining, among the above described additives, in particular, the diluent, the binder, the flavoring agent, the emulsifier, the surfactant, the solubilizer, the suspending agent, the isotonizing agent, the preservative, the antioxidant, the stabilizing agent, and the absorption promoter, as desired. Injections are formulated by combining, among the above described additives, in particular, the emulsifier, the surfactant, the solubilizer, the suspending agent, the isotonizing agent, the buffer, the preservative, the antioxidant, the stabilizing agent, and the absorption promoter, as desired.

[0079] The pharmaceutical composition in accordance with the present invention can be prepared by a well-known method. For example, to prepare tablets, a preparation process comprising steps of pre-mixing, granulating, drying, milling, main-mixing, compression, coating, and screening in this order can be applied. Either wet granulation (a non-aqueous system is preferred) or dry granulation may be employed.

[0080] In the pre-mixing step, a diluent and a binder are mixed, for example, in a 20 L super mixer. In the granulating step, to the resulting mixture, the active ingredient and an organic solvent such as ethanol are added, which are then granulated, for example, in a 20 L super mixer. In the drying step, the resulting granules are dried in a tray dryer, etc. The milling step is then conducted by a power mill, etc. To the milled granules, a disintegrant and a lubricant are added, and the main mixing step is conducted, for example, in a 10/20 L tumbler mixer, etc. Then, the compression step is conducted by a tablet press. Finally, the screening step is conducted to obtain the pharmaceutical composition (tablets).

[0081] Further, before the addition of a diluent and a binder in the pre-mixing step, another pre-mixing step wherein the active ingredient and the gelation inhibitor are added in advance can be performed. In such case, in the granulating step, only an organic solvent such as ethanol will be added. Moreover, between the coating step and the screening step, a mixing step in a 5 L tumbler mixer, etc., may be performed.

[0082] The dosage of the pharmaceutical composition in accordance with the Invention depends on symptoms, age, and dosage forms, but in general, in terms of the active ingredient, from 100 µg to 10 g thereof is administered daily once or in a few divided portions to an adult.

[0083] The pharmaceutical composition in accordance with the present invention is extremely useful as an angiogenic inhibitor, and is effective as an agent to prevent or treat diseases against which angiogenic inhibition is effective, an angiogenic inhibitor, an anti-tumor agent, an agent to treat angioma, a cancer metastasis inhibitor, an agent to treat retinal angiogenesis, an agent to treat diabetic retinopathy, an agent to treat inflammatory diseases, an agent to treat inflammatory diseases selected from the group consisting of osteoarthritis, rheumatic arthritis, psoriasis, and delayed hyperactivity, and an agent to treat atherosclerosis.

[0084] Further, when the pharmaceutical composition in accordance with the present invention is used as an anti-tumor agent, the target tumor thereof is, for example, pancreatic cancer, stomach cancer, colorectal cancer, breast cancer, prostate cancer, lung cancer, renal cancer, brain tumor, blood cancer, or ovarian cancer. In particular, stomach cancer, colorectal cancer, prostate cancer, or renal cancer is a preferred target.

[0085] Moreover, the pharmaceutical composition in accordance with the present invention exhibits a potent c-Kit kinase inhibitory action, and hence is useful as an anti-tumor agent against tumors exacerbated by activated c-Kit kinase (acute myeloid leukemia, mast cell leukemia, small cell lung cancer, GIST, testicular tumor, ovarian cancer, breast cancer, brain tumor, neuroblastoma, and colorectal cancer). The pharmaceutical composition in accordance with the present invention is also useful as an agent to treat diseases such as mastocytosis in which the involvement of c-Kit kinase is suspected, allergies, and asthma.

Examples

[0086] The present invention is further explained in detail by referring to examples and comparative examples in the following paragraphs. However, the present invention shall not be limited by the following examples by any means.

[Preparation of Medicament (Active Ingredient)]

(Preparation Example 1) Preparation (1) of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolincarboxamide

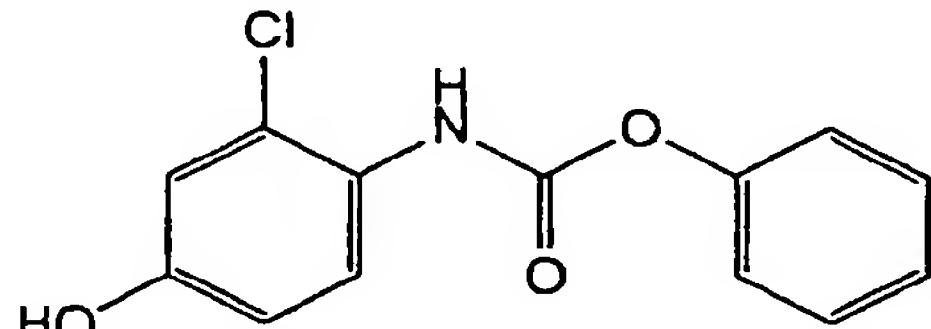
[0087] Phenyl N-(4-(6-carbamoyl-7-methoxy-4-quinolyl)oxy-2-chlorophenyl)carbamate (17.5g, 37.7 mmol) disclosed in WO 02/32872 was dissolved in *N,N*-dimethylformamide (350 mL), and then cyclopropylamine (6.53 mL, 94.25 mmol) was added to the reaction mixture under a nitrogen atmosphere, followed by stirring overnight at room temperature. To the mixture was added water (1.75L), and the mixture was stirred. Precipitated crude crystals were collected by filtration, washed with water, and dried at 70 °C for 50 min. To the obtained crude crystals was added ethanol (300 mL), and then the mixture was heated under reflux for 30 min to dissolve, followed by stirring overnight to cool slowly down to room temperature. Precipitated crystals was collected by filtration and dried under vacuum, and then further dried at 70 °C for 8 hours to give the titled crystals (12.91 g; 80.2%).

(Preparation Example 2) Preparation (2) of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolincarboxamide

(1) Preparation of phenyl N-(2-chloro-4-hydroxyphenyl)carbamate

[0088]

[Formula 3]



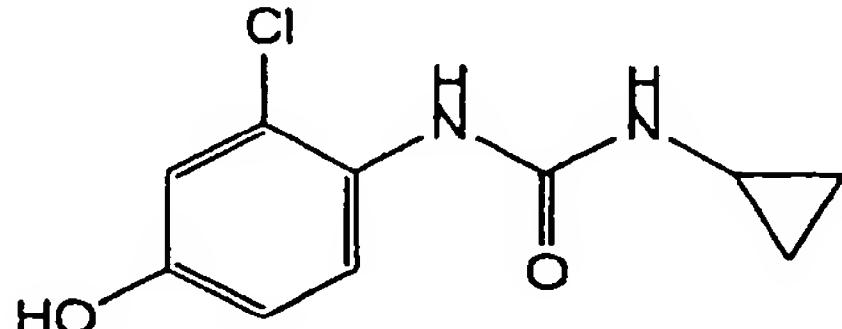
To a suspension of 4-amino-3-chlorophenol (23.7 g) in *N,N*-dimethylformamide (100 mL) was added pyridine (23.4 mL) while cooling in an ice bath, and phenyl chloroformate (23.2 mL) was added dropwise below 20 °C. After stirring at room temperature for 30 min, water (400mL), ethyl acetate (300 mL), and 6N-HCl (48 mL) were added and stirred. The organic layer was separated, washed twice with a 10% aqueous sodium chloride solution (200 mL), and dried over magnesium sulfate. The solvent was evaporated to give 46 g of the titled compound as a solid.

¹H-NMR Spectrum (CDCl₃) δ(ppm): 5.12 (1H, br s), 6.75 (1H, dd, J=9.2, 2.8 Hz), 6.92 (1H, d, J=2.8 Hz), 7.18-7.28 (4H, m), 7.37-7.43 (2H, m), 7.94 (1H, br s)

(2) Preparation of 1-(2-chloro-4-hydroxyphenyl)-3-cyclopropylurea

[0089]

[Formula 4]



To a solution of phenyl N-(2-chloro-4-hydroxyphenyl)carbamate in *N,N*-dimethylformamide (100 mL) was added cyclopropylamine (22.7 mL) while cooling in an ice bath, and the stirring was continued at room temperature overnight. Water (400 mL), ethyl acetate (300 mL), and 6N-HCl (55 mL) were added thereto, and the mixture was stirred. The organic layer was then separated, washed twice with a 10% aqueous sodium chloride solution (200 mL), and dried over magnesium sulfate. The solvent was evaporated to give prism crystals, which were collected by filtration and washed with

heptane to give 22.8 g of the titled compound (yield from 4-amino-3-chlorophenol: 77%).

¹H-NMR Spectrum (CDCl₃) δ(ppm): 0.72-0.77 (2H, m), 0.87-0.95 (2H, m), 2.60-2.65 (1H, m), 4.89 (1H, br s), 5.60 (1H, br s), 6.71 (1H, dd, J=8.8, 2.8 Hz), 6.88 (1H, d, J=2.8 Hz), 7.24-7.30 (1H, br s), 7.90 (1H, d, J=8.8 Hz)

5 (3) Preparation of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide

[0090] To dimethyl sulfoxide (20 mL) were added 7-methoxy-4-chloroquinoline-6-carboxamide (0.983 g), 1-(2-chloro-4-hydroxyphenyl)-3-cyclopropylurea (1.13 g) and cesium carbonate (2.71 g), and the mixture was heated and stirred at 10 70 °C for 23 hours. The reaction mixture was cooled to room temperature, and water (50 mL) was added, and the resultant crystals were then collected by filtration to give 1.56 g of the titled compound (yield: 88%).

(Preparation Example 3) Preparation (3) of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide

[0091] 7-Methoxy-4-chloroquinoline-6-carboxamide (5.00 kg, 21.13 mol), dimethyl sulfoxide (55.05 kg), 1-(2-chloro-4-hydroxyphenyl)-3-cyclopropylurea (5.75 kg, 25.35 mol) and potassium t-butoxide (2.85 kg, 25.35 mol) were introduced in this order into a reaction vessel under a nitrogen atmosphere. The mixture was stirred for 30 min at 20 °C, and the temperature was raised to 65 °C over 2.5 hours. The mixture was stirred at the same temperature for 19 hours. 33% (v/v) acetone-water (5.0 L) and water (10.0 L) were added dropwise over 3.5 hours. After the addition was completed, the mixture was stirred at 60 °C for 2 hours. 33% (v/v) acetone-water (20.0 L) and water (40.0 L) were added dropwise at 55 °C or more over 1 hour. After stirring at 40 °C for 16 hours, precipitated crystals were collected by filtration using a nitrogen pressure filter, and was washed with 33% (v/v) acetone-water (33.3 L), water (66.7 L), and acetone (50.0 L) in that order. The obtained crystals were dried at 60 °C for 22 hours using a conical vacuum dryer to give 7.78 kg of the titled compound (yield: 96.3%).

[0092] Further, all of the ¹H-NMR chemical shift values of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide prepared in Preparation Examples 1 to 3 described above agreed with those of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide described in WO 02/32872.

30 [Stability Evaluation of Medicament]

[0093] The crystals (C) (hereinafter, referred to as the "medicament Y") of methanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide (the "medicament X") synthesized in the above "Preparation of Medicament (Active Ingredient)" was combined with the following 10 compounds (that exhibit various pH values when 5% (w/w) aqueous solutions or suspensions were made therewith. In the table, pH values thereof are shown). Stability of the medicament X therewith was evaluated.

[0094]

[Table 1]

	pH value of 5% (w/w) aqueous solution or suspension
Magnesium oxide (MgO, Tomita Pharmaceutical Co., Ltd.)	10.63
Sodium carbonate (Na ₂ CO ₃ , Wako Pure Chemical Industries, Ltd.)	11.45
Disodium hydrogenphosphate (Na ₂ HPO ₄ , Kanto Chemical Co., Inc.)	9.26
Sodium citrate (Sodium citrate, Kozakai Pharmaceutical Co., Ltd.)	8.22
Dipotassium hydrogenphosphate (K ₂ HPO ₄ , Wako Pure Chemical Industries, Ltd.)	9.11
Sodium acetate (Sodium acetate, Wako Pure Chemical Industries, Ltd.)	8.46
Sodium hydrogencarbonate (NaHCO ₃ , Wako Pure Chemical Industries, Ltd.)	8.15

(continued)

	pH value of 5% (w/w) aqueous solution or suspension
Sodium hydroxide (NaOH, Wako Pure Chemical Industries, Ltd.)	13.56
Glycine (Glycine, Ajinomoto Co., Inc.)	6.17
δ -Gluconolactone (δ -Gluconolactone, Kanto Chemical Co., Inc.)	2.40

[0095] Anhydrous dibasic calcium phosphate (a diluent, from Kyowa Chemical Industry Co., Ltd.), croscarmellose sodium (a disintegrant, trade name: Ac-Di-Sol, from FMC International Inc.), hydroxypropylcellulose (a binder, trade name: HPC-L, from Nippon Soda Co., Ltd.), and the medicament Y are mixed in a ratio of 10/2.3/3/0.19 (w/w/w/w). To the resulting mixture, water was added, which then underwent the mixing/wet granulation process in a tablet mill followed by drying at 60°C for 5 hr. to give the pellets.

[0096] Approximately 50 mg of each of stabilizers, magnesium oxide (MgO), sodium carbonate (Na₂CO₃), disodium hydrogenphosphate (Na₂HPO₄), sodium citrate, dipotassium hydrogenphosphate (K₂HPO₄), sodium acetate, sodium hydrogencarbonate (NaHCO₃), sodium hydroxide (NaOH), glycine, and δ -gluconolactone was combined together to grind in a mortar, with which approximately 500 mg of the above pellets were kneaded in a mortar. To the resulting mixture, 50 L of water was added and mixed further.

[0097] The prepared mixture was divided into about 100 mg in 2 PP tubes, which were then stressed for a week under the conditions of 60°C/open and under the conditions of 60°C/75% relative humidity/open (hereinafter, relative humidity is abbreviated as "RH", and "open" refers to conditions wherein an open tube is heated and humidified). To the stressed mixture, 8 mL of an extractant was added, which then underwent sonication. The resulting suspension was centrifuged to give the supernatant as a sample solution, which then was analyzed by HPLC. The results are shown in Table 2. In Table 2, the results from the one with no stabilizer are also shown.

[0098]

[Table 2]

Additive	60°C/open, 1 week		60°C/75% RH/open, 1 week	
	HPLC purity (%)	Decomposed product A (%)	HPLC purity (%)	Decomposed product A (%)
No additive	97.0	0.40	95.6	1.63
MgO	97.4	0.08	97.2	0.06
Na ₂ CO ₃	97.6	0.06	97.3	0.12
Na ₂ HPO ₄	97.5	0.06	97.5	0.08
δ -Gluconolactone	97.9	0.10	95.6	1.88
Sodium citrate	97.6	0.10	97.6	0.09
K ₂ HPO ₄	97.4	0.06	97.4	0.08
Sodium acetate	97.6	0.08	97.4	0.21
Glycine	97.0	0.15	92.3	1.38
NaHCO ₃	97.5	0.11	97.3	0.10
NaOH	97.5	0.06	97.4	0.06

[0099] The relationship between the pH value of a 5% (w/w) aqueous solution or suspension of each stabilizer and the decomposed product A (see the chemical formula described above) is also shown in Figure 1. These results demonstrate that when the pH value of a 5% (w/w) aqueous solution or suspension of the stabilizer is 8 or more, the decomposition can be significantly reduced.

[Preparation of Pharmaceutical Composition]

(Example 1) 10 mg Tablets: containing magnesium oxide

5 [0100] In a 1 L super mixer 2.5 g of the medicament Y, 10 g of magnesium oxide (a stabilizer, from Tomita Pharmaceutical Co., Ltd.), 48.5 g of D-mannitol (a diluent, from Towa Chemical Industry Co., Ltd.), 10 g of partially alpha starch (a disintegrant, trade name: PCS (pharmaceutical grade), from Asahi Kasei Corporation), 22.5 g of crystalline cellulose (a diluent, trade name: Avicel PH101, from Asahi Kasei Corporation), and 3 g of hydroxypropylcellulose (a binder, trade name: HPC-L, from Nippon Soda Co., Ltd.) were mixed. Then thereto a suitable amount of purified water was added followed by granulation, drying, and milling to give the granules. To these granules, 3 g of croscarmellose sodium (a disintegrant, trade name: Ac-Di-Sol, from FMC International Inc.) and 0.5 g of magnesium stearate (a lubricant) were admixed, and then tablets were formed by a tablet press to give tablets (the total mass per tablet was 400 mg) containing 10 mg of the medicament Y per tablet.

15 (Comparative Example 1) 10 mg Tablets: containing no magnesium oxide

20 [0101] In a 1 L super mixer 2.5 g of the medicament Y, 10 g of calcium hydrogenphosphate (a diluent), 48.5 g of D-mannitol (a diluent, from Towa Chemical Industry Co., Ltd.), 10 g of partially alpha starch (a disintegrant, trade name: PCS (pharmaceutical grade), from Asahi Kasei Corporation), 22.5 g of crystalline cellulose (a diluent, trade name: Avicel PH101, from Asahi Kasei Corporation), and 3 g of hydroxypropylcellulose (a binder, trade name: HPC-L, from Nippon Soda Co., Ltd.) were mixed. Then thereto a suitable amount of purified water was added followed by granulation, drying, and milling to give the granules. To these granules, 3 g of croscarmellose sodium (a disintegrant, trade name: Ac-Di-Sol, from FMC International Inc.) and 0.5 g of magnesium stearate (a lubricant) were admixed, and then tablets were formed by a tablet press to give tablets (the total mass per tablet was 400 mg) containing 10 mg of the medicament Y per tablet.

25 [0102] Stability was tested for the tablets prepared in Example 1 and Comparative Example 1. In the test, after the tablets were stored at 5°C, at 25°C, and at 40°C and under relative humidity 75% RH, for 3 months each, impurity levels (%) were determined by HPLC. The results are shown in Table 3 below. As shown in Table 3, the tablets containing magnesium oxide (MgO) (Example 1) are superior in stability to the tablets containing no magnesium oxide (MgO) (Comparative Example 1). In particular, the stability under the humidified conditions was remarkably improved with the stabilizer.

30 [0103]

[Table 3]

Storage conditions	Example 1	Comparative 1 Example
5°C/3 Months	0	0
25°C/3 Months	0	0.17
40°C.75% RH/3 Months	0	1.58

40 values: impurity levels (%) determined by HPLC

45 [0104] Further, the ability of decomposition suppression was examined for magnesium oxide, disodium hydrogenphosphate, sodium hydrogencarbonate, and sodium hydroxide.

50 [0105] A placebo tablet containing 8.0 mg of light anhydrous silicic acid (trade name: AEROSIL (registered trademark) 200, from Nippon Aerosil Co., Ltd.), 52.5 mg of D-mannitol (from Towa Chemical Industry Co., Ltd.), 30.0 mg of crystalline cellulose (trade name: Avicel PH101, from Asahi Kasei Corporation), 3.0 mg of hydroxypropylcellulose (trade name: HPC-L, from Nippon Soda Co., Ltd.), 5.0 mg of croscarmellose sodium (trade name: Ac-Di-Sol, from FMC International Inc.), 1.5 mg of sodium stearyl fumarate (from JRS Pharma LP), and 5.0 mg of opadry yellow was prepared according to an ordinary method. Approximately 30 g of the placebo tablets were ground in a tablet mill, to which then about 33 mg of the medicament Y was added. By repetitive mixing of the medicament Y with the ground placebo tablets, 1/1000 diluted powder (0.1%) was obtained.

55 [0106] Approximately 100 mg of each of stabilizers (magnesium oxide, disodium hydrogenphosphate, sodium hydrogencarbonate, and sodium hydroxide) was mixed with 1,900 mg of the 0.1% powder in a mortar to prepare a powder containing a 5% stabilizer. Likewise, a powder containing a 4, 3, 2, or 1% stabilizer was prepared.

56 [0107] In a glass vial, approximately 200 mg of each of the prepared mixtures (0.2 mg of the medicament X is contained) was stored and stressed under the conditions of 65°C/75% RH/open for a week. To the stressed mixture, 5 mL of an

extractant was added, which then underwent sonication. The resulting suspension was centrifuged to give the supernatant as a sample solution, which then was analyzed by HPLC. The results are shown in Figure 6. Figure 6 is a graph illustrating the amount of the decomposed product A generated when various kinds of stabilizers were added at various concentrations. The results demonstrate that sodium hydroxide provided the highest stabilizing effect, the decomposition of the medicament X being reduced by adding only 1% of sodium hydroxide. Further, the stabilizing effect of magnesium oxide was similar to that of sodium hydroxide, the decomposition of the medicament X being significantly reduced by adding only 1% of magnesium oxide. The stabilizing effect of magnesium oxide was almost constant by adding 3% or more thereof.

10 [Inhibition of Gelation]

(Example 2) 1 mg Tablets

15 [0108] In a 20 L super mixer, 24 g of the medicament Y and 192 g of light anhydrous silicic acid (a gelation inhibitor, trade name: AEROSIL (registered trademark) 200, from Nippon Aerosil Co., Ltd.) were mixed, and thereto 1,236 g of D-mannitol (a diluent, from Towa Chemical Industry Co., Ltd.), 720 g of crystalline cellulose (a diluent, trade name: Avicel PH101, from Asahi Kasei Corporation), and 72 g of hydroxypropylcellulose (a binder, trade name: HPC-L, from Nippon Soda Co., Ltd.) were further added and mixed. A suitable amount of absolute ethanol was then added thereto to give the pellets containing the medicament Y. These pellets were dried in a tray dryer (at 60°C), and then the size of the pellets was controlled in a power mill to give the granules. With these granules, 120 g of croscarmellose sodium (a disintegrant, trade name: Ac-Di-Sol, from FMC International Inc.) and 36 g of sodium stearyl fumarate (a lubricant, from JRS Pharma LP) were mixed in a 20L tumbler mixer, and tablets were formed by a tablet press to give the tablets, the total mass per tablet of which was 100 mg. Further, the tablets were coated with a 10% aqueous solution of opadry yellow (OPADRY03F42069 YELLOW, from Colorcon (Japan) Limited) by a tablet coating machine to give the coated tablets, the total mass per tablet of which was 105 mg.

(Example 3) 10 mg Tablets

30 [0109] In a 20 L super mixer, 60 g of the medicament Y and 192 g of light anhydrous silicic acid (a gelation inhibitor, trade name: AEROSIL (registered trademark) 200, from Nippon Aerosil Co., Ltd.) were mixed, and thereto 1,200 g of D-mannitol (a diluent, from Towa Chemical Industry Co., Ltd.), 720 g of crystalline cellulose (a diluent, trade name: Avicel PH101, from Asahi Kasei Corporation), and 72 g of hydroxypropylcellulose (a binder, trade name: HPC-L, from Nippon Soda Co., Ltd.) were further added and mixed. A suitable amount of absolute ethanol was then added thereto to give the pellets containing the medicament Y. These pellets were dried in a tray dryer (at 60°C), and then the size of the pellets was controlled in a power mill to give the granules. With these granules, 120 g of croscarmellose sodium (a disintegrant, trade name: Ac-Di-Sol, from FMC International Inc.) and 36 g of sodium stearyl fumarate (a lubricant, from JRS Pharma LP) were mixed in a 20L tumbler mixer, and tablets were formed by a tablet press to give the tablets, the total mass per tablet of which was 400 mg. Further, the tablets were coated with a 10% aqueous solution of opadry yellow (OPADRY03F42069 YELLOW, from Colorcon (Japan) Limited) by a tablet coating machine to give the coated tablets, the total mass per tablet of which was 411 mg.

(Example 4) 100 mg Tablets

45 [0110] In a 1 L super mixer, 31.4 g of the medicament Y and 4 g of light anhydrous silicic acid (a gelation inhibitor, trade name: AEROSIL (registered trademark) 200, from Nippon Aerosil Co., Ltd.) were mixed, and thereto 40.1 g of anhydrous dibasic calcium phosphate (a diluent, from Kyowa Chemical Industry Co., Ltd.), 10 g of low-substituted hydroxypropylcellulose (a binder, trade name: L-HPC (LH-21), from Shin-Etsu Chemical Co., Ltd.), and 3 g of hydroxypropylcellulose (a binder, trade name: HPC-L, from Nippon Soda Co., Ltd.) were further added and mixed. A suitable amount of absolute ethanol was then added thereto to give the pellets containing the medicament Y. These pellets were dried in a tray dryer (at 60°C), and then the size of the pellets was controlled in a power mill to give the granules. With these granules, 10 g of croscarmellose sodium (a disintegrant, trade name: Ac-Di-Sol, from FMC International Inc.) and 1.5 g of sodium stearyl fumarate (a lubricant, from JRS Pharma LP) were mixed, and tablets were formed by a tablet press to give the tablets, the total mass per tablet of which was 400 mg.

55 (Comparative Example 2) 100 mg Tablets

[0111] In a 1 L super mixer, 31.4 g of the medicament Y, 44.1 g of anhydrous dibasic calcium phosphate (a diluent, from Kyowa Chemical Industry Co., Ltd.), 10 g of low-substituted hydroxypropylcellulose (a binder, trade name: L-HPC

(LH-21), from Shin-Etsu Chemical Co., Ltd.), and 3 g of hydroxypropylcellulose (a binder, trade name: HPC-L, from Nippon Soda Co., Ltd.) were mixed. A suitable amount of absolute ethanol was then added thereto to give the pellets containing the medicament Y. These pellets were dried in a tray dryer (at 60°C), and then the size of the pellets was controlled in a power mill to give the granules. With these granules, 10 g of croscarmellose sodium (a disintegrant, trade name: Ac-Di-Sol, from FMC International Inc.) and 1.5 g of sodium stearyl fumarate (a lubricant, from JRS Pharma LP) were mixed in a tumbler mixer, and tablets were formed by a tablet press to give the tablets, the total mass per tablet of which was 400 mg.

(Example 5) 100 mg Tablets: 8% of light anhydrous silicic acid

[0112] In a 1 L super mixer, 31.4 g of the medicament Y and 8 g of light anhydrous silicic acid (a gelation inhibitor, trade name: AEROSIL (registered trademark) 200, from Nippon Aerosil Co., Ltd.) were mixed, and thereto 42.1 g of anhydrous dibasic calcium phosphate (a diluent, from Kyowa Chemical Industry Co., Ltd.), and 10 g of low-substituted hydroxypropylcellulose (a binder, trade name: L-HPC (LH-21), from Shin-Etsu Chemical Co., Ltd.) were further added and mixed. A suitable amount of absolute ethanol with 2 g of hydroxypropylcellulose (a binder, trade name: HPC-L, from Nippon Soda Co., Ltd.) suspended therein was then added thereto to give the pellets containing the medicament of the present invention. These pellets were dried in a tray dryer (at 60°C), and then the size of the pellets was controlled in a power mill to give the granules. With these granules, 5 g of croscarmellose sodium (a disintegrant, trade name: Ac-Di-Sol, from FMC International Inc.) and 1.5 g of sodium stearyl fumarate (a lubricant, from JRS Pharma LP) were mixed, and tablets were formed by a tablet press to give the tablets, the total mass per tablet of which was 400 mg.

(Example 6) 100 mg Tablets: 6% of light anhydrous silicic acid

[0113] In a 1 L super mixer, 31.4 g of the medicament Y and 6 g of light anhydrous silicic acid (a gelation inhibitor, trade name: AEROSIL (registered trademark) 200, from Nippon Aerosil Co., Ltd.) were mixed, and thereto 44.1 g of anhydrous dibasic calcium phosphate (a diluent, from Kyowa Chemical Industry Co., Ltd.), and 10 g of low-substituted hydroxypropylcellulose (a binder, trade name: L-HPC (LH-21), from Shin-Etsu Chemical Co., Ltd.) were further added and mixed. A suitable amount of absolute ethanol with 2 g of hydroxypropylcellulose (a binder, trade name: HPC-L, from Nippon Soda Co., Ltd.) suspended therein was then added thereto to give the pellets containing the medicament of the present invention. These pellets were dried in a tray dryer (at 60°C), and then the size of the pellets was controlled in a power mill to give the granules. With these granules, 5 g of croscarmellose sodium (a disintegrant, trade name: Ac-Di-Sol, from FMC International Inc.) and 1.5 g of sodium stearyl fumarate (a lubricant, from JRS Pharma LP) were mixed, and tablets were formed by a tablet press to give the tablets, the total mass per tablet of which was 400 mg.

(Example 7) 100 mg Tablets: 4% of light anhydrous silicic acid

[0114] In a 1 L super mixer, 31.4 g of the medicament Y and 4 g of light anhydrous silicic acid (a gelation inhibitor, trade name: AEROSIL (registered trademark) 200, from Nippon Aerosil Co., Ltd.) were mixed, and thereto 46.1 g of anhydrous dibasic calcium phosphate (a diluent, from Kyowa Chemical Industry Co., Ltd.), and 10 g of low-substituted hydroxypropylcellulose (a binder, trade name: L-HPC (LH-21), from Shin-Etsu Chemical Co., Ltd.) were further added and mixed. A suitable amount of absolute ethanol with 2 g of hydroxypropylcellulose (a binder, trade name: HPC-L, from Nippon Soda Co., Ltd.) suspended therein was then added thereto to give the pellets containing the medicament of the present invention. These pellets were dried in a tray dryer (at 60°C), and then the size of the pellets was controlled in a power mill to give the granules. With these granules, 5 g of croscarmellose sodium (a disintegrant, trade name: Ac-Di-Sol, from FMC International Inc.) and 1.5 g of sodium stearyl fumarate (a lubricant, from JRS Pharma LP) were mixed, and tablets were formed by a tablet press to give the tablets, the total mass per tablet of which was 400 mg.

[0115] The storage test and the dissolution test were conducted for the tablets prepared above according to the methods described below.

(Storage Test)

[0116] Tablets in a glass bottle with the cap open were stored at 5°C, at 60°C and 75% RH, at 40°C and 75% RH, or at 30°C and 65% RH.

(Dissolution test)

[0117] The dissolution test was conducted according to the Japanese Pharmacopoeia 14th Edition and by the paddle method under the conditions described below. The test solution: 900 mL of 0.1 mol/L hydrochloric acid. The rotation

speeds: 50 rpm. The temperature of the test solution: 37°C.

(Experimental Example 1)

[0118] The storage test (the storage duration was 3 months) was conducted for the tablets prepared in Examples 2 and 3. No delayed dissolution was found for the tablets of both Examples at any of the conditions at 5°C, at 30°C and 65% RH, and at 40°C and 75% RH. The results of each of the examples are shown in Figures 2 and 3.

(Experimental Example 2)

[0119] The tablets prepared in Example 4 and Comparative Example 2 were stored at 60°C and 75% RH for 7 days. The tablets then underwent the dissolution test. The results are shown in Figure 4. For the tablets of Comparative Example 2, the gelation of the tablet surface was noted even at the beginning of the test. Further, significant delayed dissolution was observed after the storage test. On the other hand, for the tablets of Example 4, the gelation on the surface was not noted on any of the tablets. The inhibition of delayed dissolution after the storage was also observed.

(Experimental Example 3)

[0120] The tablets prepared in Examples 5 to 7 were stored at 60°C and 75% RH for 7 days. The tablets then underwent the dissolution test. To confirm the influence of the amount of light anhydrous silicic acid contained in the tablet, the drug released (%) 30 min. after the dissolution test started was compared. The results are shown in Figure 5. When from 16 to 32 mg of light anhydrous silicic acid was contained per 100 mg of the medicament Y, delayed dissolution seen in Comparative Example 2 was not observed. In particular, for the tablets of Example 5, wherein 32 mg of light anhydrous silicic acid was contained, delayed dissolution was barely seen after the storage as well.

[0121] The results shown above demonstrate that the addition of from 4 to 8% of the gelation inhibitor provides the pharmaceutical composition comprising the medicament X with great dissolution properties, while the gelation was effectively inhibited. Next examined were disintegration properties when higher levels of the gelation inhibitor was contained in the pharmaceutical composition. Disintegration properties when the stabilizer and the gelation inhibitor were contained therein were also examined. Disintegration properties were further examined, when silicone dioxide hydrate or calcium silicate was used in place of light anhydrous silicic acid as a gelation inhibitor.

(Comparative Example 3) 25 mg Tablets

[0122] To 7.85 g of the medicament Y, 22.4 g of D-mannitol (a diluent, from Towa Chemical Industry Co., Ltd.), 15.0 g of crystalline cellulose (a diluent, trade name: Avicel PH101, from Asahi Kasei Corporation), and 1.5 g of hydroxypropylcellulose (a binder, trade name: HPC-L, from Nippon Soda Co., Ltd.) were added and mixed. A suitable amount of absolute ethanol was then added thereto to give the pellets containing the medicament Y. These pellets were dried in a tray dryer (at 60°C), and then the size of the pellets was controlled in a small speed mill to give the granules. With these granules, 2.5 g of croscarmellose sodium (a disintegrant, trade name: Ac-Di-Sol, from FMC International Inc.) and 0.8 g of sodium stearyl fumarate (a lubricant, from JRS Pharma LP) were mixed, and tablets were formed by a tablet press to give the tablets, the total mass per tablet of which was 200 mg.

(Example 8) 25 mg Tablets: 12% of light anhydrous silicic acid

[0123] In a 1 L super mixer, 7.85 g of the medicament Y and 6 g of light anhydrous silicic acid (a gelation inhibitor, trade name: AEROSIL (registered trademark) 200, from Nippon Aerosil Co., Ltd.) were mixed, and thereto 16.4 g of D-mannitol (a diluent, from Towa Chemical Industry Co., Ltd.), 15.0 g of crystalline cellulose (a diluent, trade name: Avicel PH101, from Asahi Kasei Corporation), and 1.5 g of hydroxypropylcellulose (a binder, trade name: HPC-L, from Nippon Soda Co., Ltd.) were further added and mixed. A suitable amount of absolute ethanol was then added thereto to give the pellets containing the medicament Y. These pellets were dried in a tray dryer (at 60°C), and then the size of the pellets was controlled in a small speed mill to give the granules. With these granules, 2.5 g of croscarmellose sodium (a disintegrant, trade name: Ac-Di-Sol, from FMC International Inc.) and 0.8 g of sodium stearyl fumarate (a lubricant, from JRS Pharma LP) were mixed, and tablets were formed by a tablet press to give the tablets, the total mass per tablet of which was 200 mg.

(Example 9) 25 mg Tablets: 20% of light anhydrous silicic acid

[0124] In a 1 L super mixer, 7.85 g of the medicament Y and 10 g of light anhydrous silicic acid (a gelation inhibitor,

trade name: AEROSIL (registered trademark) 200, from Nippon Aerosil Co., Ltd.) were mixed, and thereto 12.4 g of D-mannitol (a diluent, from Towa Chemical Industry Co., Ltd.), 15.0 g of crystalline cellulose (a diluent, trade name: Avicel PH101, from Asahi Kasei Corporation), and 1.5 g of hydroxypropylcellulose (a binder, trade name: HPC-L, from Nippon Soda Co., Ltd.) were further added and mixed. A suitable amount of absolute ethanol was then added thereto to give the pellets containing the medicament Y. These pellets were dried in a tray dryer (at 60°C), and then the size of the pellets was controlled in a small speed mill to give the granules. With these granules, 2.5 g of croscarmellose sodium (a disintegrant, trade name: Ac-Di-Sol, from FMC International Inc.) and 0.8 g of sodium stearyl fumarate (a lubricant, from JRS Pharma LP) were mixed, and tablets were formed by a tablet press to give the tablets, the total mass per tablet of which was 200 mg.

(Example 10) 25 mg Tablets: 8% of light anhydrous silicic acid and 3% magnesium oxide

[0125] In a 1 L super mixer, 15.7 g of the medicament Y and 8 g of light anhydrous silicic acid (a gelation inhibitor, trade name: AEROSIL (registered trademark) 200, from Nippon Aerosil Co., Ltd.) were mixed, and thereto 3 g of magnesium oxide (a stabilizer, from Tomita Pharmaceutical Co., Ltd.), 33.8 g of D-mannitol (a diluent, from Towa Chemical Industry Co., Ltd.), 30 g of crystalline cellulose (a diluent, trade name: Avicel PH101, from Asahi Kasei Corporation), and 3 g of hydroxypropylcellulose (a binder, trade name: HPC-L, from Nippon Soda Co., Ltd.) were further added and mixed. A suitable amount of absolute ethanol was then added thereto to give the pellets containing the medicament Y. These pellets were dried in a tray dryer (at 60°C), and then the size of the pellets was controlled in a small speed mill to give the granules. With these granules, 5 g of croscarmellose sodium (a disintegrant, trade name: Ac-Di-Sol, from FMC International Inc.) and 1.5 g of sodium stearyl fumarate (a lubricant, from JRS Pharma LP) were mixed, and tablets were formed by a tablet press to give the tablets, the total mass per tablet of which was 200 mg.

(Example 11) 25 mg Tablets: 8% of light anhydrous silicic acid and 5% disodium hydrogenphosphate

[0126] In a 1 L super mixer, 7.85 g of the medicament Y and 4 g of light anhydrous silicic acid (a gelation inhibitor, trade name: AEROSIL (registered trademark) 200, from Nippon Aerosil Co., Ltd.) were mixed, and thereto 2.5 g of disodium hydrogenphosphate (a stabilizer, from Kanto Chemical Co., Inc.), 15.9 g of D-mannitol (a diluent, from Towa Chemical Industry Co., Ltd.), 15 g of crystalline cellulose (a diluent, trade name: Avicel PH101, from Asahi Kasei Corporation), and 1.5 g of hydroxypropylcellulose (a binder, trade name: HPC-L, from Nippon Soda Co., Ltd.) were further added and mixed. A suitable amount of absolute ethanol was then added thereto to give the pellets containing the medicament Y. These pellets were dried in a tray dryer (at 60°C), and then the size of the pellets was controlled in a small speed mill to give the granules. With these granules, 2.5 g of croscarmellose sodium (a disintegrant, trade name: Ac-Di-Sol, from FMC International Inc.) and 0.8 g of sodium stearyl fumarate (a lubricant, from JRS Pharma LP) were mixed, and tablets were formed by a tablet press to give the tablets, the total mass per tablet of which was 200 mg.

(Example 12) 25 mg Tablets: 8% of silicon dioxide hydrate

[0127] In a 1 L super mixer, 7.85 g of the medicament Y and 4 g of silicon dioxide hydrate (a gelation inhibitor, trade name: Sylsia, from Fuji Sylsia Chemical Ltd.) were mixed, and thereto 18.4 g of D-mannitol (a diluent, from Towa Chemical Industry Co., Ltd.), 15.0 g of crystalline cellulose (a diluent, trade name: Avicel PH101, from Asahi Kasei Corporation), and 1.5 g of hydroxypropylcellulose (a binder, trade name: HPC-L, from Nippon Soda Co., Ltd.) were further added and mixed. A suitable amount of absolute ethanol was then added thereto to give the pellets containing the medicament Y. These pellets were dried in a tray dryer (at 60°C), and then the size of the pellets was controlled in a small speed mill to give the granules. With these granules, 2.5 g of croscarmellose sodium (a disintegrant, trade name: Ac-Di-Sol, from FMC International Inc.) and 0.8 g of sodium stearyl fumarate (a lubricant, from JRS Pharma LP) were mixed, and tablets were formed by a tablet press to give the tablets, the total mass per tablet of which was 200 mg.

(Example 13) 25 mg Tablets: 8% of calcium silicate

[0128] In a 1 L super mixer, 7.85 g of the medicament Y and 4 g of calcium silicate (a gelation inhibitor, trade name: Florite (registered trademark), from Tokuyama Corp.) were mixed, and thereto 18.4 g of D-mannitol (a diluent, from Towa Chemical Industry Co., Ltd.), 15.0 g of crystalline cellulose (a diluent, trade name: Avicel PH101, from Asahi Kasei Corporation), and 1.5 g of hydroxypropylcellulose (a binder, trade name: HPC-L, from Nippon Soda Co., Ltd.) were further added and mixed. A suitable amount of absolute ethanol was then added thereto to give the pellets containing the medicament Y. These pellets were dried in a tray dryer (at 60°C), and then the size of the pellets was controlled in a small speed mill to give the granules. With these granules, 2.5 g of croscarmellose sodium (a disintegrant, trade name: Ac-Di-Sol, from FMC International Inc.) and 0.8 g of sodium stearyl fumarate (a lubricant, from JRS Pharma LP) were

mixed, and tablets were formed by a tablet press to give the tablets, the total mass per tablet of which was 200 mg. [0129] The disintegration test was conducted for the tablets prepared above by the method described in the Japanese Pharmacopoeia 14th Edition. The disintegration time is summarized in Table 4.

[0130]

5

[Table 4]

Sample	Disintegration time
Comparative Example 3	15 min. or longer
Example 8	1.2 to 1.4 min.
Example 9	0.9 to 1.1 min.
Example 10	3.9 to 4.1 min.
Example 11	2.9 to 3.1 min.
Example 12	7.6 to 8.2 min.
Example 13	2.3 to 2.5 min.

[0131] The tablets of any of Examples 8 to 13 had a shorter disintegration time than those from Comparative Example 3. It is shown that disintegration properties of the tablets prepared in Examples 8 to 13 were superior. The above demonstration confirms that the pharmaceutical composition of the present invention inhibits the gelation effectively.

(Formulation Examples)

25

[0132] Formulation examples comprising the crystals (C) of methanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide (the medicament X) synthesized in the above "Preparation of Medicament (Active Ingredient)" are shown. In Table 5, a formulation of a 10 mg tablet (a coated tablet) and in Table 6, a formulation of a 100 mg tablet (a coated tablet) are illustrated.

[0133]

[Table 5]

Material	Purpose	Amount contained (mg)
The compound	Active ingredient	12.3
Magnesium oxide	Stabilizer	10
Anhydrous dibasic calcium phosphate	Diluent	150.7
D-Mannitol	Diluent	153
Partially alpha starch	Disintegrant	20
Crystalline cellulose	Diluent.	16
Hydroxypropylcellulose	Binder	12
Subtotal		374
Croscarmellose sodium	Disintegrant	20
Sodium stearyl fumarate	Lubricant	6
Subtotal		400
Opadry yellow	Coating agent	11
Total		411

55 [0134]

[Table 6]

Material	Purpose	Amount contained (mg)
The compound	Active ingredient	122.5
Magnesium oxide	Stabilizer	10
Anhydrous dibasic calcium phosphate	Diluent	37.5
Partially alpha starch	Disintegrant	20
Croscarmellose sodium	Disintegrant	20
Purified water	Solvent	q.s.
Subtotal		210
Anhydrous dibasic calcium phosphate	Disintegrant	136
Croscarmellose sodium	Lubricant	8
Crystalline cellulose	Diluent	16
Hydroxypropylcellulose	Disintegrant	4
Purified water	Solvent	q.s.
Subtotal		374
Croscarmellose sodium	Disintegrant	20
Sodium stearyl fumarate	Lubricant	6
Subtotal		400
Opadry yellow	Coating agent	11
Total		411

[0135] In Tables 5 and 6, "the compound" refers to the crystals (C) of methanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide (the medicament X), and "opadry yellow" refers to a pre-mixed materials consisting of Hydroxypropylmethylcellulose 2910, talc, Macrogol 6000 (molecular weight: 8,000), titanium oxide, and yellow iron sesquioxide in 56.0, 28.0, 10.0, 4.0, and 2.0% (w/w), respectively.

[0136] A 10 mg tablet was formulated by the following processes. The active ingredient, magnesium oxide, anhydrous dibasic calcium phosphate, D-mannitol, partially alpha starch, crystalline cellulose, and hydroxypropylcellulose were mixed, and then thereto a suitable amount of purified water was added to prepare pellets. These pellets were dried, and then the size of the pellets was controlled. To the resulting granules, croscarmellose sodium and sodium stearyl fumarate were added and mixed, and a tablet was formed. On the resulting tablet, a film of opadry yellow was coated by a fluidized bed coating technique.

[0137] A 100 mg tablet was formulated by the following processes. The active ingredient, magnesium oxide, anhydrous dibasic calcium phosphate, partially alpha starch, and croscarmellose sodium were mixed, and then thereto a suitable amount of purified water was added to prepare pellets. These pellets were dried, and then the size of the pellets was controlled. To the resulting pellets, anhydrous dibasic calcium phosphate, croscarmellose sodium, crystalline cellulose, and hydroxypropylcellulose were added and mixed, and then thereto a suitable amount of purified water was added to prepare granules. These granules were dried, and then the size of the granules was controlled. To the resulting granules, croscarmellose sodium and sodium stearyl fumarate were added and mixed, and a tablet was formed. On the resulting tablet, a film of opadry yellow was coated by a fluidized bed coating technique.

50 Industrial Applicability

[0138] The pharmaceutical composition in accordance with the present invention is highly stable, and hence is clinically useful.

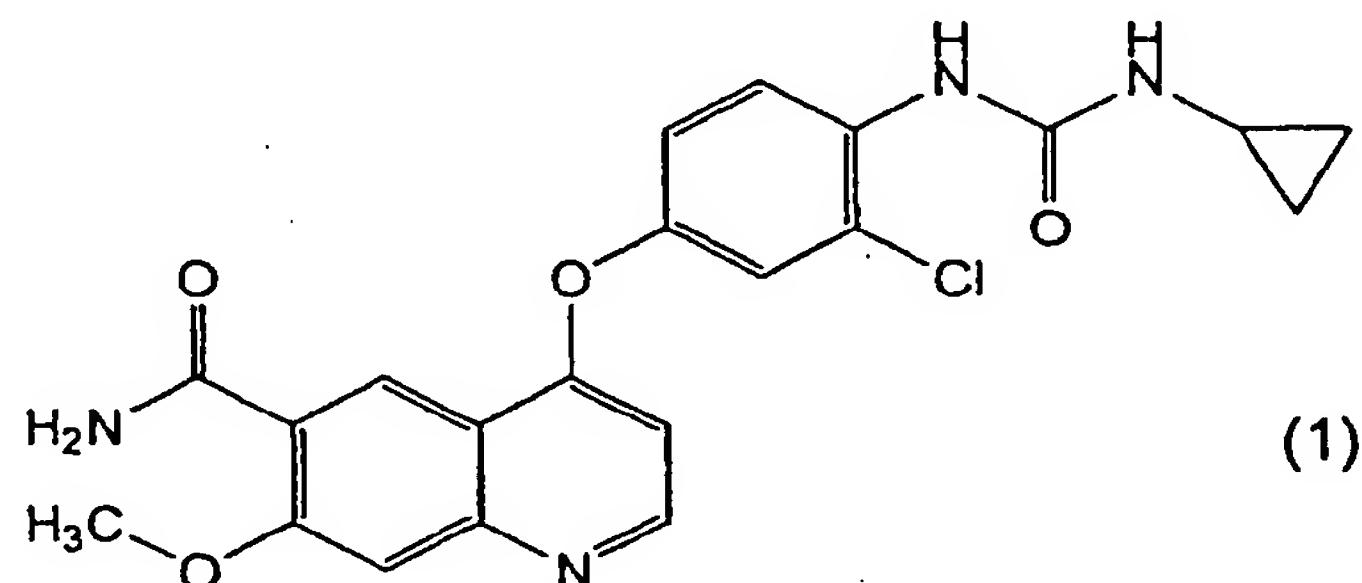
55 Claims

1. A pharmaceutical composition comprising:

an active ingredient consisting of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide represented by Formula (1) described below, salt thereof, or solvate of the foregoing; and

5 (i) a compound, a 5% (w/w) aqueous solution or suspension of which has a pH of 8 or more, and/or
 (ii) silicic acid, salt thereof, or solvate of the foregoing.

10 [Formula 1]



2. The pharmaceutical composition according to Claim 1, wherein (i) the compound, a 5% (w/w) aqueous solution or suspension of which has a pH of 8 or more, is one or more selected from the group consisting of magnesium oxide, calcium oxide, sodium carbonate, disodium hydrogenphosphate, sodium citrate, dipotassium hydrogenphosphate, sodium acetate, sodium hydrogencarbonate, and sodium hydroxide.
3. The pharmaceutical composition according to Claim 1 or Claim 2, wherein (ii) silicic acid, salt thereof, or solvate of the foregoing is one or more selected from the group consisting of light anhydrous silicic acid, silicon dioxide hydrate, calcium silicate, magnesium silicate, magnesium aluminosilicate, magnesium aluminometasilicate, magnesium aluminum silicate, synthetic aluminum silicate, and hydrous silicic dioxide.
4. The pharmaceutical composition according to any one of Claims 1 to 3, wherein the active ingredient is crystals of hydrochloride, hydrobromide, p-toluenesulfonate, sulfate, methanesulfonate, or ethanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide or crystals of solvate thereof.
5. The pharmaceutical composition according to any one of Claims 1 to 3, wherein the active ingredient is crystals of methanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide or crystals of solvate thereof.
6. The pharmaceutical composition according to any one of Claims 1 to 3, wherein the active ingredient is crystals of methanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide.
7. The pharmaceutical composition according to any one of Claims 1 to 3, wherein the active ingredient is crystals of hydrate of methanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide.
8. The pharmaceutical composition according to any one of Claims 1 to 3, wherein the active ingredient is crystals of dimethyl sulfoxide solvate of methanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide.
9. The pharmaceutical composition according to any one of Claims 1 to 3, wherein the active ingredient is crystals of acetic acid solvate of methanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide.
10. The pharmaceutical composition according to Claim 6, wherein the crystals of methanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide are crystals (A) having diffraction peaks at diffraction angles ($2\theta \pm 0.2^\circ$) of 9.65° and 18.37° in a powder X-ray diffraction.

11. The pharmaceutical composition according to Claim 6, wherein the crystals of methanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide are crystals (B) having diffraction peaks at diffraction angles ($2\theta \pm 0.2^\circ$) of 5.72° and 13.84° in a powder X-ray diffraction.

5 12. The pharmaceutical composition according to Claim 6, wherein the crystals of methanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide are crystals (C) having diffraction peaks at diffraction angles ($2\theta \pm 0.2^\circ$) of 14.20° and 17.59° in a powder X-ray diffraction.

10 13. The pharmaceutical composition according to Claim 7, wherein the crystals of hydrate of methanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide are crystals (F) having diffraction peaks at diffraction angles ($2\theta \pm 0.2^\circ$) of 8.02° and 18.14° in a powder X-ray diffraction.

15 14. The pharmaceutical composition according to Claim 9, wherein the crystals of acetic acid solvate of methanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide are crystals (I) having diffraction peaks at diffraction angles ($2\theta \pm 0.2^\circ$) of 9.36° and 12.40° in a powder X-ray diffraction.

20 15. The pharmaceutical composition according to Claim 10, wherein the crystals (A) having diffraction peaks at diffraction angles ($2\theta \pm 0.2^\circ$) of 9.65° and 18.37° in a powder X-ray diffraction are prepared by a preparing process comprising mixing 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide, a solvent, and methanesulfonic acid to dissolve.

25 16. The pharmaceutical composition according to Claim 10, wherein the crystals (A) having diffraction peaks at diffraction angles ($2\theta \pm 0.2^\circ$) of 9.65° and 18.37° in a powder X-ray diffraction are prepared by a preparing process comprising mixing 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide, acetic acid, and methanesulfonic acid to dissolve.

30 17. The pharmaceutical composition according to Claim 11, wherein the crystals (B) having diffraction peaks at diffraction angles ($2\theta \pm 0.2^\circ$) of 5.72° and 13.84° in a powder X-ray diffraction are prepared by a preparing process comprising drying crystals (I) of acetic acid solvate of methanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide to remove acetic acid.

35 18. The pharmaceutical composition according to Claim 12, wherein the crystals (C) having diffraction peaks at diffraction angles ($2\theta \pm 0.2^\circ$) of 14.20° and 17.59° in a powder X-ray diffraction are prepared by a preparing process comprising heating crystals of dimethyl sulfoxide solvate of methanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide.

40 19. The pharmaceutical composition according to Claim 12, wherein the crystals (C) having diffraction peaks at diffraction angles ($2\theta \pm 0.2^\circ$) of 14.20° and 17.59° in a powder X-ray diffraction are prepared by a preparing process comprising mixing crystals (I) of acetic acid solvate of methanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide and a solvent.

45 20. The pharmaceutical composition according to Claim 12, wherein the crystals (C) having diffraction peaks at diffraction angles ($2\theta \pm 0.2^\circ$) of 14.20° and 17.59° in a powder X-ray diffraction are prepared by a preparing process comprising mixing 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide, acetic acid, and methanesulfonic acid to dissolve.

50 21. The pharmaceutical composition according to Claim 12, wherein the crystals (C) having diffraction peaks at diffraction angles ($2\theta \pm 0.2^\circ$) of 14.20° and 17.59° in a powder X-ray diffraction are prepared by a preparing process comprising humidifying crystals (B) of methanesulfonate of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide.

55 22. The pharmaceutical composition according to Claim 13, wherein the crystals (F) having diffraction peaks at diffraction angles ($2\theta \pm 0.2^\circ$) of 8.02° and 18.14° in a powder X-ray diffraction are prepared by a preparing process comprising mixing 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide, acetic acid, and methanesulfonic acid to dissolve.

23. The pharmaceutical composition according to Claim 14, wherein the crystals (I) having diffraction peaks at diffraction angles ($2\theta \pm 0.2^\circ$) of 9.36° and 12.40° in a powder X-ray diffraction are prepared by a preparing process comprising

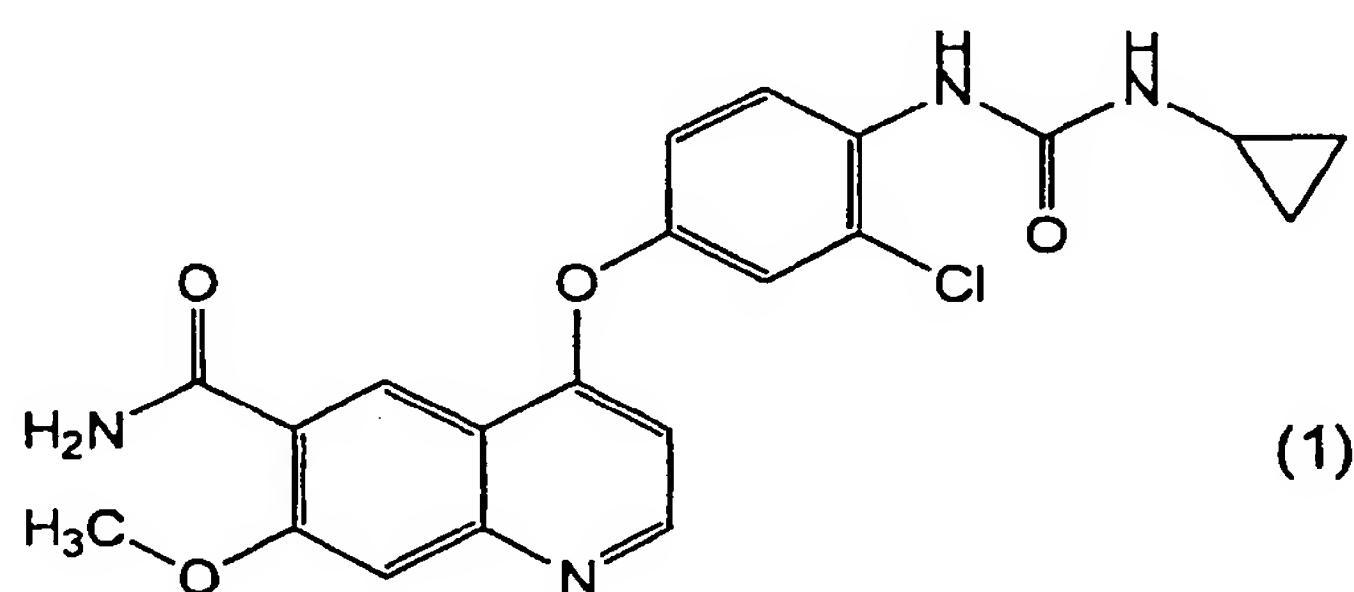
mixing 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide, acetic acid, and methanesulfonic acid to dissolve.

5 24. A process for improving stability of a pharmaceutical composition comprising an active ingredient consisting of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide represented by Formula (1) described below, salt thereof, or solvate of the foregoing, comprising adding a compound, a 5% (w/w) aqueous solution or suspension of which has a pH of 8 or more.

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[Formula 2]

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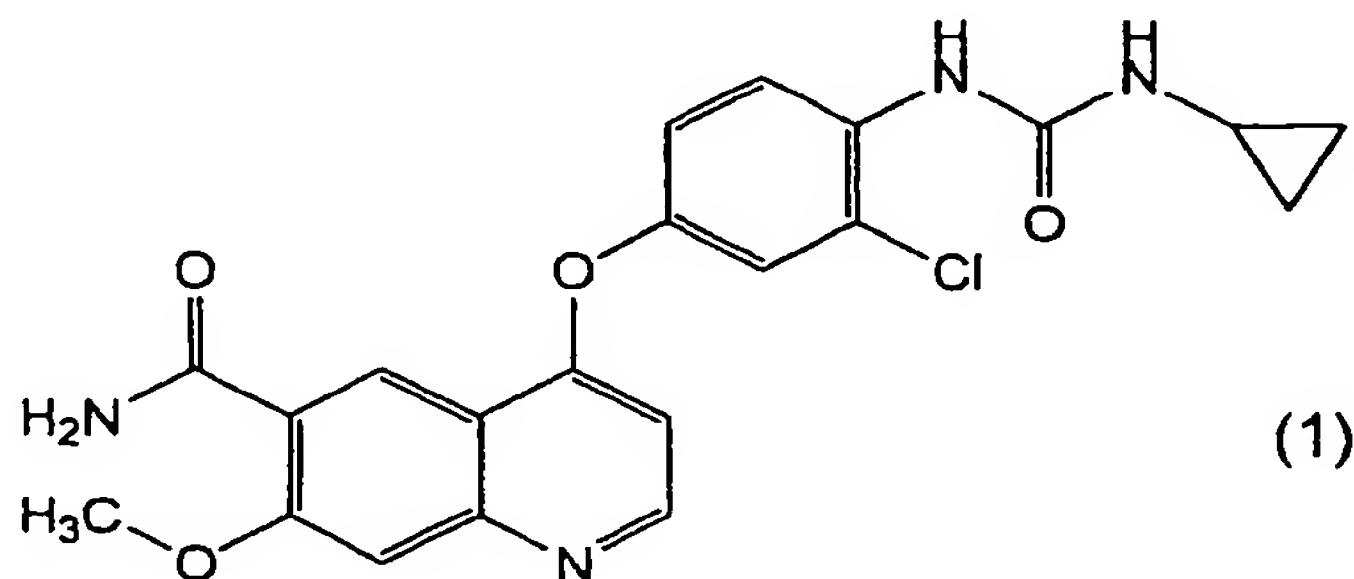
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30 25. A process for inhibiting gelation of a pharmaceutical composition comprising an active ingredient consisting of 4-(3-chloro-4-(cyclopropylaminocarbonyl)aminophenoxy)-7-methoxy-6-quinolinecarboxamide represented by Formula (1) described below, salt thereof, or solvate of the foregoing, comprising adding silicic acid, salt thereof, or solvate of the foregoing.

35

[Formula 3]

40



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Fig. 1

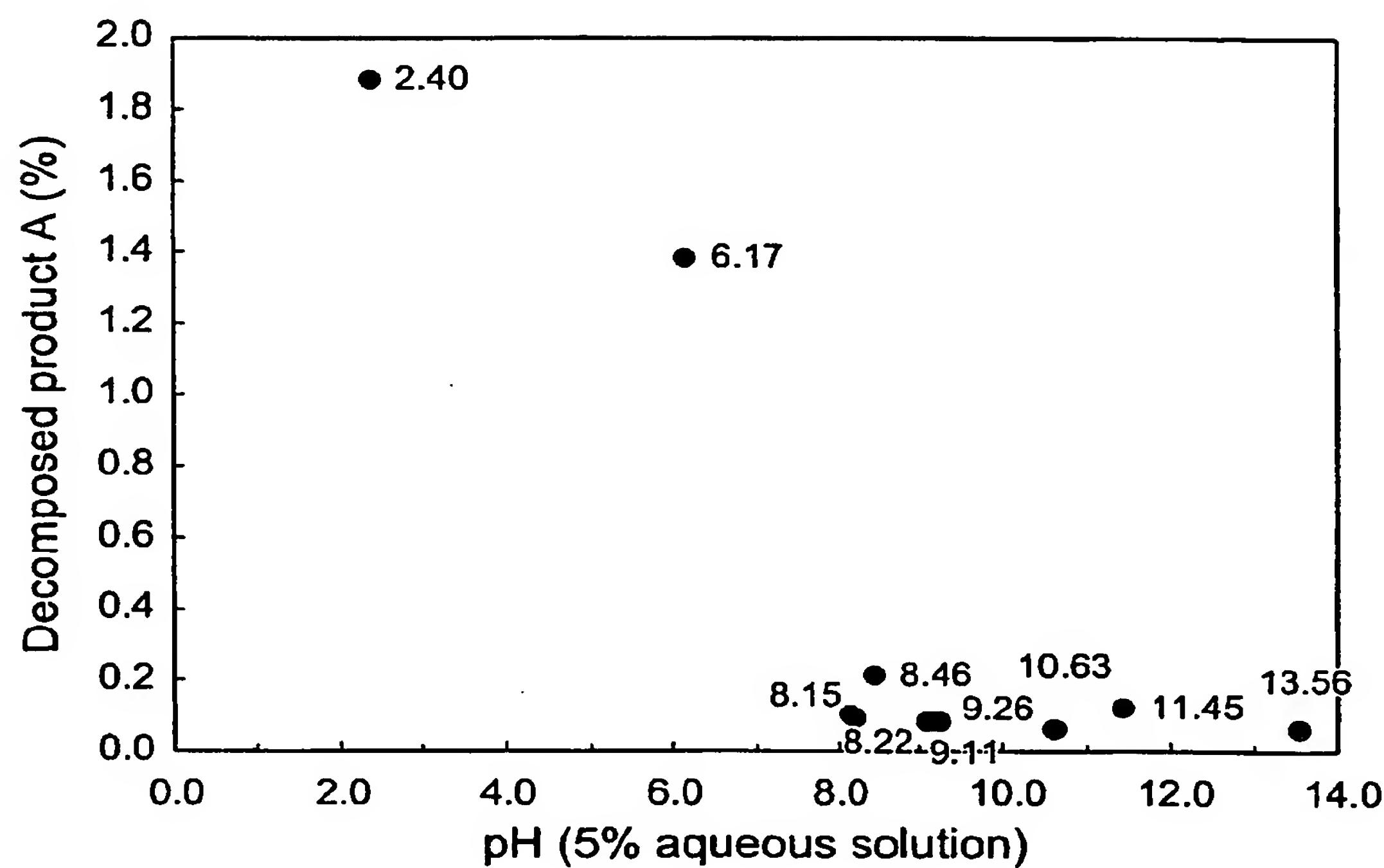


Fig.2

Dissolution test results for Example 2

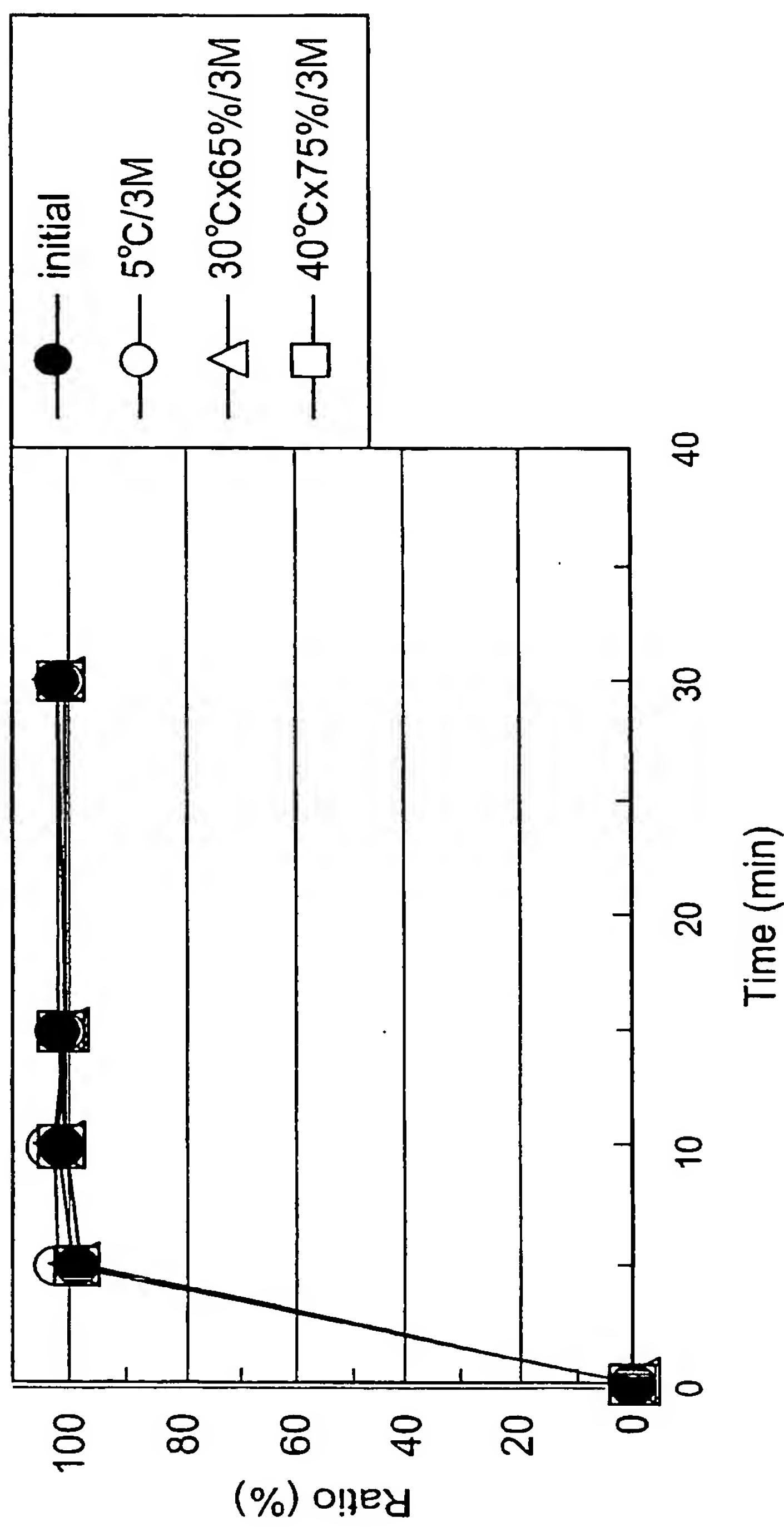


Fig.3

Dissolution test results for tablets of Example 3

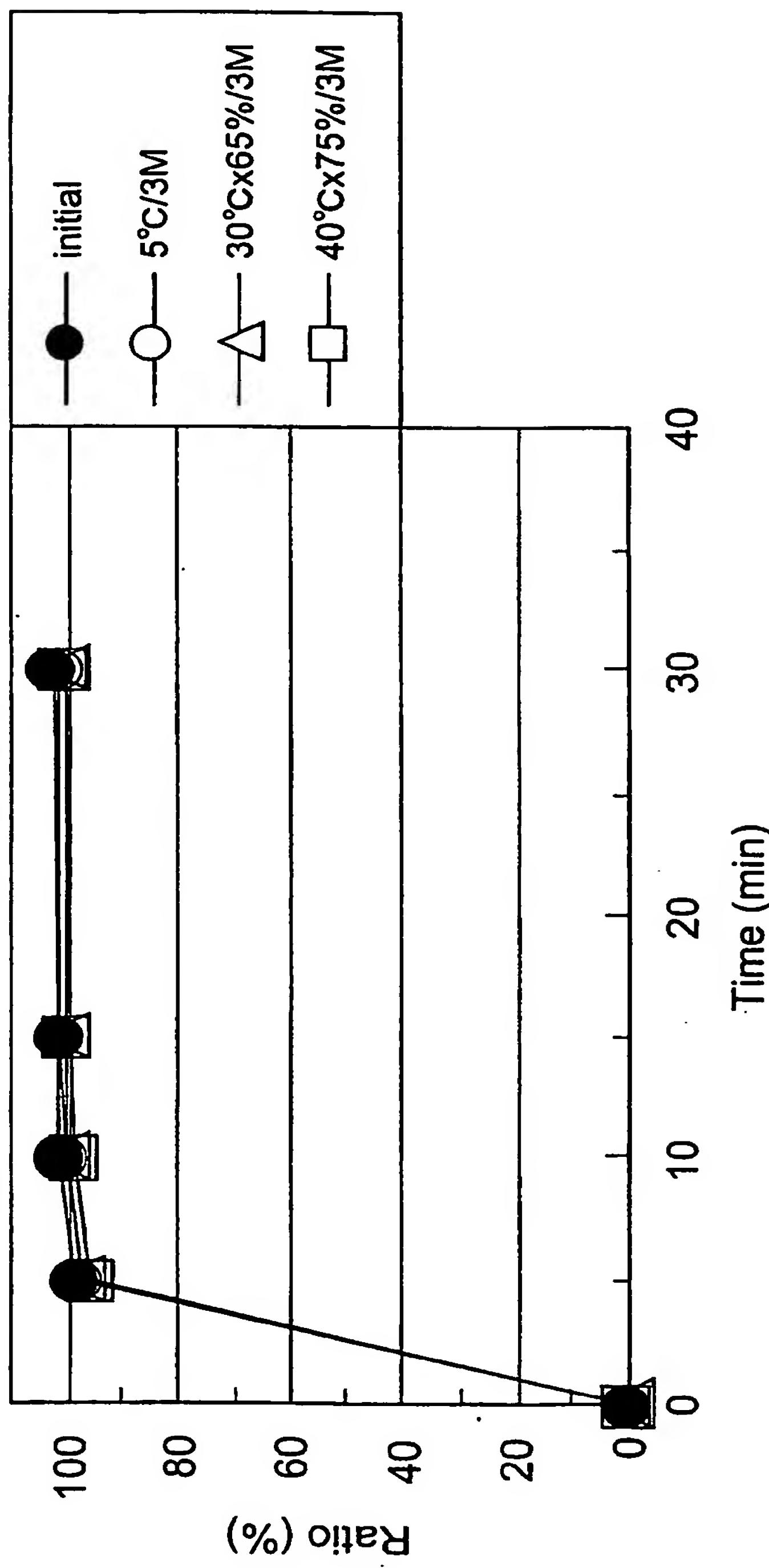


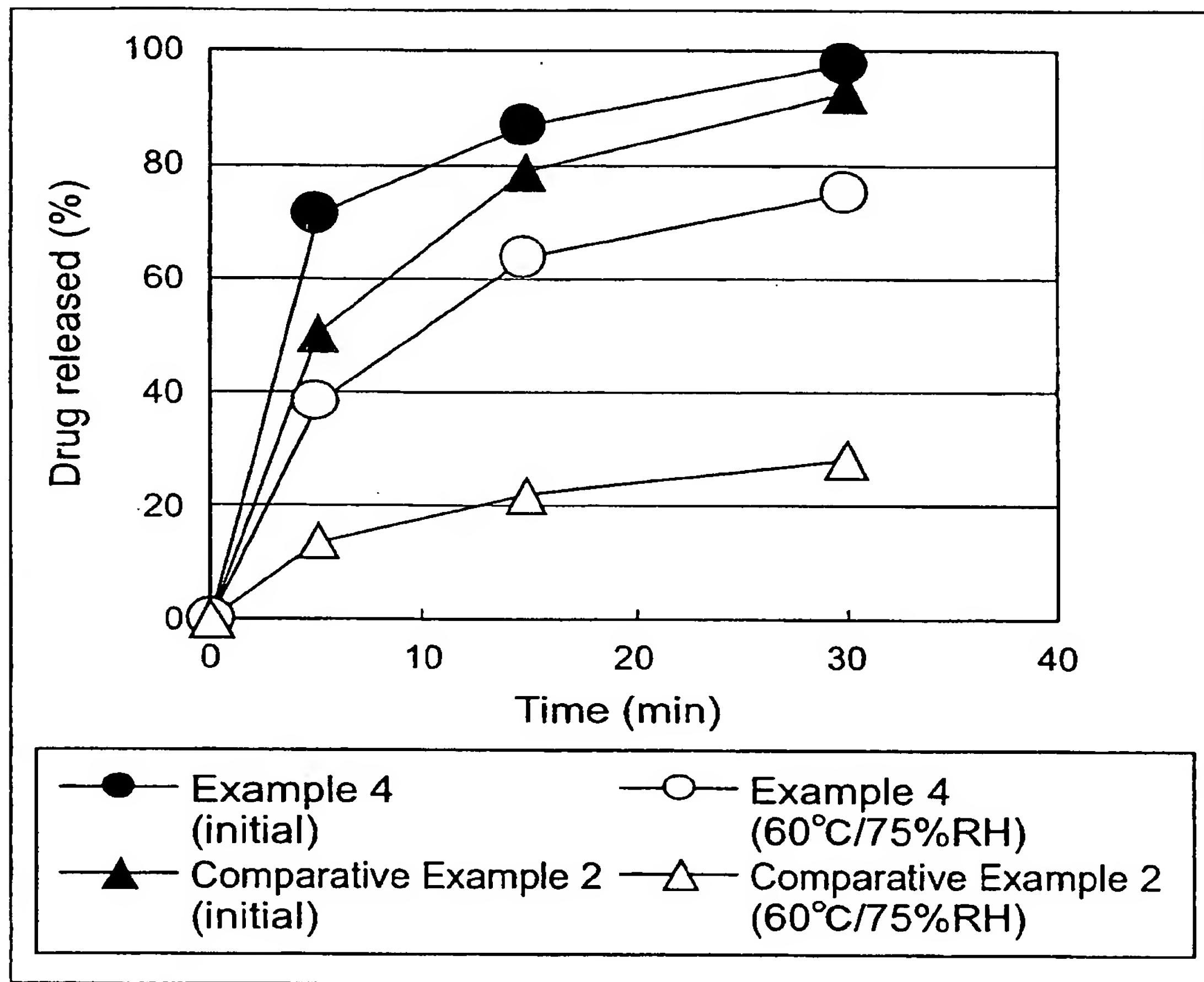
Fig.4

Fig.5

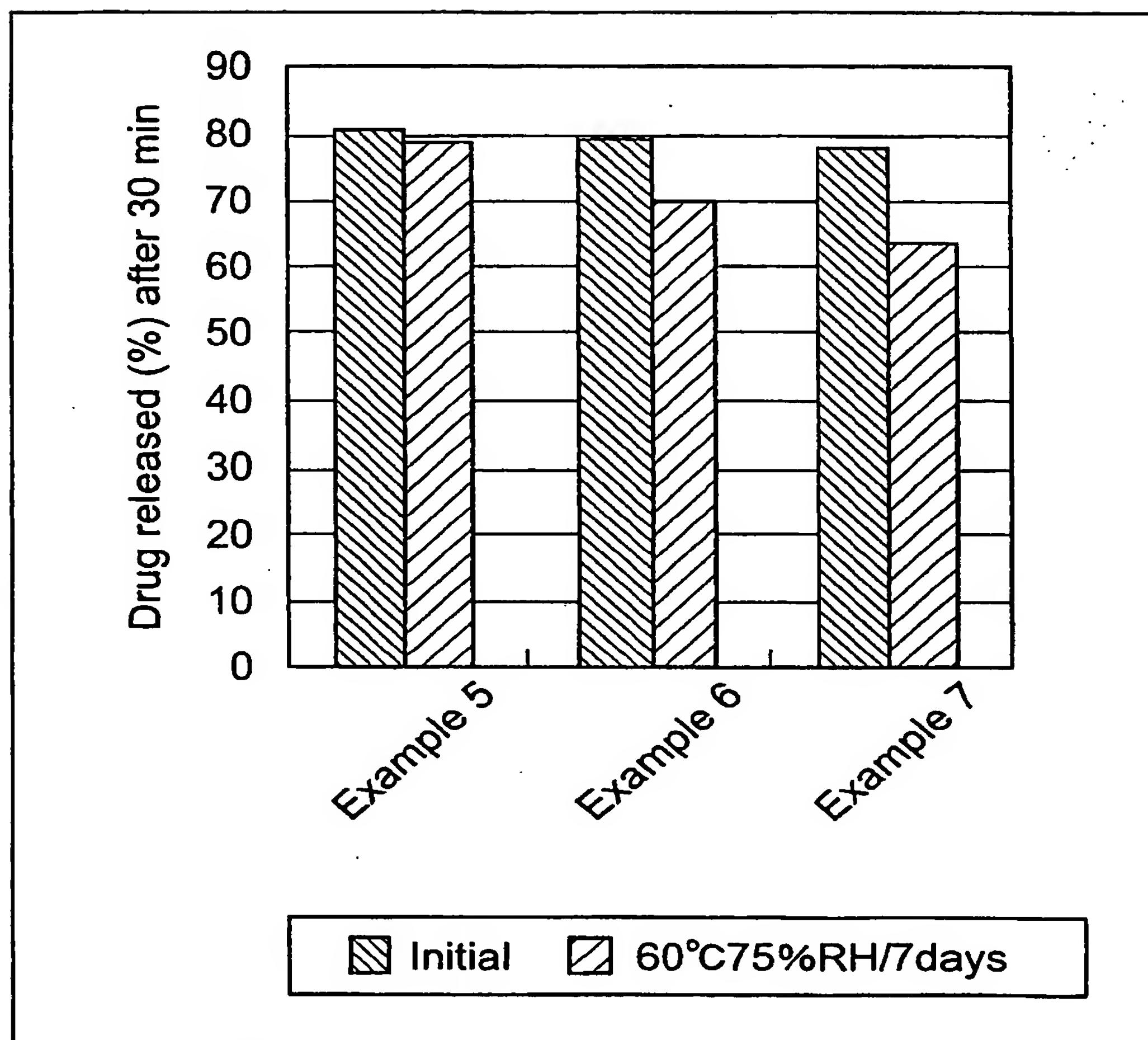
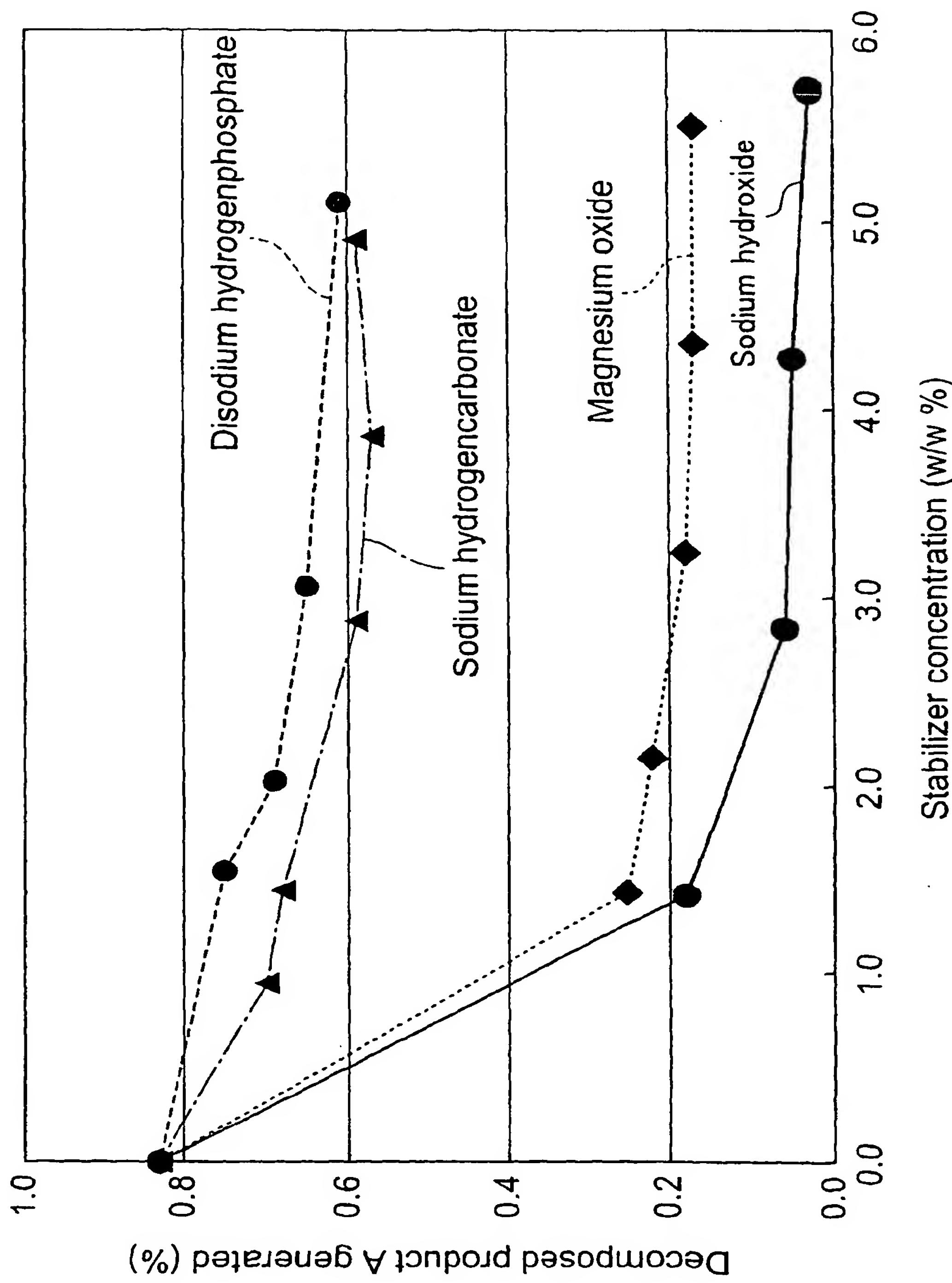


Fig. 6



INTERNATIONAL SEARCH REPORT

International application No.

PCT/JP2005/016941

A. CLASSIFICATION OF SUBJECT MATTER

A61K31/47 (2006.01), **A61K47/04** (2006.01), **A61P9/00** (2006.01), **A61P35/00** (2006.01), **A61P43/00** (2006.01), **C07D215/48** (2006.01)

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

A61K31/47 (2006.01), **A61K47/04** (2006.01), **A61P9/00** (2006.01), **A61P35/00** (2006.01), **A61P43/00** (2006.01), **C07D215/48** (2006.01)

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Jitsuyo Shinan Koho	1922-1996	Jitsuyo Shinan Toroku Koho	1996-2005
Kokai Jitsuyo Shinan Koho	1971-2005	Toroku Jitsuyo Shinan Koho	1994-2005

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
CAplus (STN), REGISTRY (STN)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	WO 2002/032872 A1 (Eisai Co., Ltd.), 25 April, 2002 (25.04.02), Claim 1; page 74, line 33 to page 75, line 2; example 368 & EP 1415987 A1 & US 2004/53908 A1	1-25
Y	JP 11-501343 A (Buckman Laboratories International, Inc.), 02 February, 1999 (02.02.99), Claims 1, 17 & WO 96/26997 A1 & EP 822971 A1	1-23, 25
Y	JP 63-28427 A (Eisai Co., Ltd.), 06 February, 1988 (06.02.88), Claim 1; page 2, upper left column, lines 1 to 16 (Family: none)	1-23, 25

 Further documents are listed in the continuation of Box C. See patent family annex.

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Date of the actual completion of the international search 01 November, 2005 (01.11.05)	Date of mailing of the international search report 15 November, 2005 (15.11.05)
Name and mailing address of the ISA/ Japanese Patent Office Facsimile No.	Authorized officer Telephone No.

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INTERNATIONAL SEARCH REPORT

International application No.

PCT/JP2005/016941

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	WO 00/71097 A1 (Takeda Chemical Industries, Ltd.), 30 November, 2000 (30.11.00), Claim 1; page 6, line 29 to page 7, line 13 & JP 2001-31567 A & AU 4778500 A	1-24
Y	JP 2004-155773 A (Takeda Chemical Industries, Ltd.), 03 June, 2004 (03.06.04), Claims 1 to 26 & WO 2004/35052 A1	1-24
Y	JP 2003-26576 A (Eisai Co., Ltd.), 29 January, 2003 (29.01.03), Par. No. [0021] (Family: none)	1-25
Y	JP 2002-3365 A (Eisai Co., Ltd.), 09 January, 2002 (09.01.02), Par. No. [0018] (Family: none)	1-25

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REFERENCES CITED IN THE DESCRIPTION

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Patent documents cited in the description

- WO 0232872 A [0003] [0028] [0087] [0092]